第六十九 麻藥條約

關スル條約廠薬ノ製造制限及分配取締ニ

一九三五年(昭和十年)四月十七日批准一九三二年(昭和六年)十月十三日、ジェネーグ」。 炊予署名

一九三五年(昭和十年)六月三日伊催音客託一九三五年(昭和十年)四月十七日批准

一九三五年(昭和十年)大月十二日公布

一九三五年(昭和十年)九月一日帝國ニ對シ效力移生

第一章 定義(第一條)

第二章 見待(第二條乃至第五條)

第三章 製造制限(第六條乃至第九條)

统四章 禁止及制限(第十條乃至第十二條)

防穴章 行政規定(第十五條乃至第十九條)第五章 取締(第十三條及第十四條)

第七章 一般規定(第二十條乃至第三十四條)

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ティン」共和國大統領、壊地利共和國聯邦大統領、白支 獨逸國大統領、「アメリカ」合衆國大統領、「アルゼン

NO. 69. STUPÉFIANTS.

CONVENTION FOR LIMITING THE MANU-FACTURE AND REGULATING THE DIS-TRIBUTION OF NARCOTIC DRUGS.

Signed at Geneva, July 13, 1931.

Ratified April 17, 1935.

Ratification deposited June 3, 1935:

Promulgated June 12, 1935.

Effective in respect of Japan from September 1,

SUMMARY.

CHAPTER I. DEFINITIONS (Art. 1).

CHAPTER II. ESTIMATES (Art. 2 to 5).

CHAPTER III. LIMITATION OF MANUFACTURE (Art. 6 to 9).

CHAPTER IV. PROHIBITIONS AND RESTRICTIONS (Art. 10 to 12).

CHAPTER V. CONTROL (Arts. 13 and 14).

CHAPTER VI. ADMINISTRATIVE PROVISIONS (Art. 15 to 19).

CHAPTER VII. GENERAL PROVISIONS (Art. 20 to 34).

THE PRESIDENT OF THE GERMAN REICH; THE PRESIDENT OF THE UNITED STATES OF AMERICA;

入八八五

耳義國皇帝陛下、「ボリヴィア」 北和國大統領、「ブラ ジル」合衆共和國大統領、「グレート、ブリテン」、 「アイルランド」及「グレート、プリテン」海外領土皇 帝印度皇帝陛下、「チリ」北和國大統領、「コスタ、リ カ」共和國大統領、「キュバ」共和國大統領、丁採國及 「アイスランド」國皇帝陛下、「ダンチッヒ」自由市ノ 為ニ「ボーランド」共和國大統領、「ドミニカ」共和國 大統領、「エジプト」國皇帝陛下、西班牙共和國假政府 大統領、「エティオピア」國皇帝陛下、佛蘭西共和國大 統領、希臘共和國大統領、「グァティラ」共和國大統 領、「(デァーズ」及「ネデド」例就二個地皇帝陛下、伊 太利國皇帝陸下、日本國皇帝陛下、「リベリア」共和國 大統領、「リスアニア」、共和國大統領、「ルクセンブル グ」因大公殿下、「メキシコ」合衆國大統領、「モナコ」 **岡公殿下、「バナァ」共和國大統領、「バラグァイ」共和** 岡大統領、和蘭國皇帝陛下、「ベルシァ」國皇帝陛下、 「ボーランド」共和國大統領、「ボルトガル」共和國大 統領、「ルーマニア」國皇帝陛下、「サン、マリノ」共和 岡攝政官、選雜國皇帝陛下、瑞典國皇帝陛下、瑞西聯 邦政府、「チェッコスロヴァキア」北和國大統領、「ウル

THE PRESIDENT OF THE ARGENTINE REPUBLIC; THE FEDERAL PRESIDENT OF THE AUSTRIAN REPUBLIC; HIS MAJESTY THE KING OF THE BELGIANS; THE PRESIDENT OF THE REPUBLIC OF BOLIVIA; THE PRESIDENT OF THE REPUBLIC OF THE UNITED STATES OF BRAZIL; HIS MAJESTY THE KING OF GREAT BRITAIN, IRELAND AND THE BRITISH DOMINIONS BEYOND THE SEAS, EMPEROR OF INDIA; THE PRESI-DENT OF THE REPUBLIC OF CHILE; THE PRESIDENT OF THE REPUBLIC OF COSTA RICA; THE PRESIDENT OF THE REPUBLIC OF CUBA; HIS MAJESTY THE KING OF DENMARK AND ICELAND; THE PRESIDENT OF THE POLISH REPUBLIC, FOR THE FREE CVIY OF DANZIG; THE PRESIDENT OF THE DOMINICAN RE-PUBLIC; HIS MAJESTY THE KING OF EGYPT; THE PRESIDENT OF THE PROVISIONAL GOVERNMENT OF THE SPANISH REPUBLIC; HIS MAJESTY THE EM-PEROR AND KING OF THE KINGS OF ABYSSINIA; THE PRESIDENT OF THE FRENCH REPUBLIC; THE PRESI-DENT OF THE HELLENIC REPUBLIC; THE PRESIDENT OF THE REPUBLIC OF GUATEMALA; HIS MAJESTY THE KING OF HEJAZ, NEJD AND DEPENDENCIES; HIS MAJESTY THE KING OF ITALY; HIS MAJESTY THE EMPEROR OF JAPAN; THE PRESIDENT OF THE RE-PUBLIC OF LIBERIA; THE PRESIDENT OF THE RE-PUBLIC OF LITHUANIA; HER ROYAL HIGHNESS THE GRAND DUCHESS OF LUXEMBURG; THE PRESIDENT OF THE UNITED STATES OF MEXICO; HIS SERENE HIGHNESS THE PRINCE OF MONACO; THE PRESIDENT

グァイ」 共和國大統領、「ヴェネズエラ」合衆國大統領

現定ヲ補足センコトヲ欲シ「ジュネーヴ」1於テ署名セラレタル 國際阿片條約ノ「ジュネーヴ」1於テ署名セラレタル 國際阿片條約ノ三日「ヘーグ」1於ラ及 干九百二十五年二月十九日並ニ其ノ分配ヲ取締リ 以テ 干九百十二年一月二十需要ニ制限スルコトヲ國際協定ニ依り有效ナラシメ麻薬ノ製造ヲ醫療用及學術用ノ為ノ世界ノ正當ナル

委員ヲ任命セリ之ガ爲條約ヲ締結スルコトニ決シ左ノ如ク其ノ全權

獨逸國大統領

インパーベン」 大官(体職)「ケェルネル、フライヘル、フォン、ラ OF THE REPUBLIC OF PANAMA; THE PRESIDENT OF THE REPUBLIC OF PARAGUAY; HER MAJESTY THE QUEEN OF THE NETHERLANDS; HIS IMPERIAL MAJESTY THE SHAH OF PERSIA; THE PRESIDENT OF THE POLISH REPUBLIC; THE PRESIDENT OF THE PORTUGUESE REPUBLIC; HIS MAJESTY THE KING OF ROUMANIA; I CAPITANI REGGENTI OF THE REPUBLIC OF SAN MARINO; HIS MAJESTY THE KING OF SIAM; HIS MAJESTY THE KING OF SIAM; HIS MAJESTY THE KING OF SWEDEN; THE SWISS FEDERAL COUNCIL; THE PRESIDENT OF THE CZECHOSLOVAK REPUBLIC; THE PRESIDENT OF THE REPUBLIC OF URUGUAY; THE PRESIDENT OF THE UNITED STATES OF VENEZUELA,

Desiring to supplement the provisions of the International Opium Conventions, signed at The Hague on January 28rd, 1912, and at Geneva on February 19th, 1925, by rendering effective by in ernational agreement the limitation of the manufacture of narcotic drugs to the world's legitimate requirements for medical and scientific purposes and by regulating their distribution,

Have resolved to conclude a Convention for that purpose and have appointed as their Plenipotentiaries:

The President of the German Reich:

M. Werner Freiherr von Rheinbaben, "Staatssekretür z.D."; **國逸國内終省參事官「ドクトル、フルディール、** カーンシー

「アメリカ」合衆國大統領

國務省「ジャン、ケー、コーグドウェグ」

麻薬部長「ハリー、ジェー、アンスリンガー」

合衆國公衆衛生部精神衛生課長、醫 務 總 監 補 「ウォルター、ルーイス、トレッドウェイ」

「カリフォルニア」州上院議員「サンポーン、ヤン LA

「アルゼンティン」共和國大統領

伊太利國駐翁特命全機大使「ドクトル、フェルナ ソディイフドー

墺地利共和國聯邦大統領

國際聯盟ニ派造ノ常任代表者、特命全權公使「ニ メリシスプラリューググー

阿片及他ノ危险藥品ノ取引ニ關スル諸問委員合 委員、警察部長、宮中參事官「ドクトル、ブル ノー、ツィグシー

Dr. Waldemar KAHLER, Ministerial Counsellor at the Ministry of Interior of the Reich.

The President of the United States of America:

Mr. John K. Caldwell, of the Department of State:

Mr. Harry J. Anslinger, Commissioner of Narcotics:

Mr. Walter Lewis TREADWAY, M.D., F.A.C.P., Assistant Surgeon-General, United States Public Health, Service Chief, Division of Mental Hygiene;

Mr. Sanborn Young, Member of the Senate of the State of California.

The President of the Argentine Republic:

Dr. Fernando Perez, Ambassador Extraordinary and Plenipotentiary to His Majesty the King of Italy.

The Federal President of the Austrian Republic:

M. Emerich Pflügl, Envoy Extraordinary and

Minister Plenipotentiary, Permanent Representative accredited to the League of Nations;
Dr. Bruno Schultz, Police Director and "Conseiller aulique", Member of the Advisory Committee on Traffic in Opium and Other Dangerous Drugs.

白耳義國皇帝陛下

在「ハル」薬局依在長官「ドクトル、エフ、ド、ミッ トヤーラー

「ボリヴィア」共和國大統領

阿片及他ノ危险藥品ノ取引ニ闢スル諮問委員合 委員「ドクトル、エメ、クェリァール」

「プラジル」合衆共和國大統領

瑞西聯邦駐箚特命全権公使「ラウル、ド、リオ、ブ רע לעו

「グレート・ブリテン」、「アイルランド」及「グレート、 ブリテン」、海外領土皇帝印度皇帝陛下

「グレート、ブリテン」及北部「アイルランド」並ニ 國際聯盟ノ傾個ノ聯盟國ニ非ザル英帝國ノ一切

内務行常任文官補「サー、マルコム、デレヴィン 11 11 1

「セナダ」

恩給及國民保健省麻藥部長、大佐「シー、エイチ、

His Majesty the King of Belgium:

Dr. F. DE MYTTENAERE, Principal Inspector of Chemistry at Hal.

The President of the Republic of Bolivia:

Dr. M. CUELLAR, Member of the Advisory Committee on Traffic in Opium and Other Dangerous Drugs.

The President of the Republic of the United States of Brazil:

M. Raul do Rio Branco, Envoy Extraordinary and Minister Plenipotentiary to the Swiss Federal Council.

His Majesty the King of Great Britain, Ireland and the British Dominions beyond the Seas, Emperor of India:

For Great Britain and Northern Ireland and all parts of the British Empire which are not separate Members of the League of

Sir Malcolm Delevingne, K.C.B., Permanent Deputy-Under-Secretary in the Home Office.

For the Dominion of Canada:

Colonel C. H. L. SHARMAN, C.M.G., C.B.E.,

Hデ、ツィートソ7

ルター、エー、リデル」
國際聯盟ニ派遣ノ「カナダ」顧問「ドクトル、ウォ

印度

ビュー印度委員會委員「ドクトル、アール、ビー、バラニ

「チリ」共和國大統領

ルド」國際聯盟ニ派遣ノ常設代表部員「エンリケ、ガハ

グェレド、ロラー在「ジュネーヴ」領事「ドクトル、ヴィリアト、フィ「コスタ、リカ」共和國大統領

「キュパ」共和國大統領

「ギリュルモ、デ、プランク」國際聯盟ニ派遣ノ常任代表委員、特命全權公使

しゃケトグ、ムングミンノアシメン スト

ン」 在「ベルヌ」 代理公使「グスターフ、ラスムッセ丁扶國及「アイスランド」國皇帝陛下 Chief Narcotic Division, Department Pensions and National Health;

Dr. Walter A. RIDDELL, M.A., Ph.D., Dominion of Canada Advisory Officer accredited to the League of Nations.

For India:

Dr. R. P. PARANJPYE, Member of the Council of India.

The President of the Republic of Chile:

M. Enrique Gajardo, Member of the Permanent Delegation accredited to the League of Nations.

The President of the Republic of Costa Rica:
Dr. Viriato Figueredo Lora, Consul at Geneva.

The President of the Republic of Cuba.

M. Guillermo DE BLANCK, Envoy Extraordinary and Minister Plenipotentiary, Permanent Delegate accredited to the League of Nations; Dr. Benjamin PRIMELLES.

His Majesty the King of Denmark and Iceland:
M. Gustav Rasmussen, Chargé d'affaires at
Berne.

領「ダンチッセ」自由市ノ為ニ「ボーランド」共和國大統

ソソア、ソカル」 国際聯盟 1 派遣ノ常任代表委員、全権な使「フラ

「ドミニカ」共和國大統領

「ティー、ダブリュー、ラッセル、パシァ」「カイロ」警察部長衆麻藥ニ關スル中央情報局長「エジプト」國皇帝陛下

作「ジュキーシ」熱簡単「ツァググ、アッケグトン」

西班牙共和國假政府大統領

「エティオピア」國皇帝陛下

ト」な、伯符「ラガルド」国際聯盟ニ派遣ノ代表者、全権企使、「エントラ

佛蘭西國領事「ガストン、ブルゴア」佛蘭西共和國大統領

The President of the Polish Republic (for the Free City of Danzig):

M. François Sokal, Minister Plenipotentiary, Permanent Delegate accredited to the League of Nations.

The President of the Dominican Republic:

M. Charles Ackermann, Consul-General at Geneva.

His Majesty the King of Egypt:

T. W. Russell Pasha, Chief of Police of Cairo and Director of the Central Bureau for Information with regard to Narcotics.

The President of the Provisional Government of the Spanish Republic:

M. Julio Casares, Head of Section at the Ministry for Foreign Affairs.

His Majesty the Emperor and King of the Kings of Abyssinia:

Count Lagarde, Duke of Entotto, Minister Plenipotentiary, Representative accredited to the League of Nations.

The President of the French Republic:

M. Gaston Bourgois, Consul of France.

H ディットートッ7

ルター、エー、リデル」 國際聯盟ニ派遣ノ「カナダ」顧問「ドクトル、ウォ

印度

ピー」印度委員會委員「ドクトル、アール、ピー、バラニ

「チリ」共和國大統領

ルド」國際聯盟ニ派遣ノ常設代表部員「エンリケヽガハ

「コスタ、リカ」共和國大統領

グェレド、ロラ」在「ジュネーツ」領事「ドクトル、ツィリアト、フィ

「キュバ」共和國大統領

「ギリェルモ、ディヴランク」国際聯盟ニ派遣ノ常任代表委員、特命全権公使

しドクトグノムシングリメリ スト

丁抹國及「アイスランド」國皇帝陸下

た「ベルヌ」 代理公使「グスターフ、ラスムッセ

Chief Narcotic Division, Department Pensions and National Health:

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For India:

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領「ダンチッと」自由市ノ為ニ「ボーランド」共和國大統

ンソア、ソカルー国際聯盟ニ派遣ノ常任代表委員、全権企使「フラ

「ドミニカ」共和國大統領

在「ジュキーグ」熱阻率「シァググ、アッケグトン」

「エジプト」國皇帝陛下

「ティー、ダブリュー、ラッセル、バシァ」「カイロ」警察部長乗麻薬ニ開スル中央情報局長

西班牙共和國假政府大統領

外務省課長「フリオ、カサレス」

「エティオピア」國皇帝陛下

ト」公、伯符「ラガルド」 國際聯盟ニ派遣ノ代表者、全権企使、「エントラ

佛蘭西共和國大統領

佛蘭西國領事「ガストン、ブグゴア」

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. The President of the Dominican Republic:

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Count Lagarde, Duke of Entotto, Minister Plenipotentiary, Representative accredited to the League of Nations.

The President of the French Republic:

M. Gaston Bourgois, Consul of France.

希臘共和國大統領

エル」国際聯盟ニ派遣ノ常任代表委員「アール、ラファ

「グァテァラ」共和國大統領

ネス、モント」國立中等學校質驗心理學教師「ルイス、マルティ

「ヘデァーズ」及「ネデド」國立二國地皇帝陛下

ズ」 英國駐箚特命を權公使「シェイク、ハフィズ、ワー

伊大利國皇帝建下

ニー 前券働大臣、上院議員「ステファノ、カヴァッツォ

日本國皇帝陛下

國際聯盟帝國事務局長、全權公使澤田節藏

行政課長、內務書記官大達茂雄

「リベリア」共和國大統領

The President of the Hellenic Republic:

M. R. Raphael, Permanent Delegate accredited to the League of Nations.

The President of the Republic of Guatemala:

M. Luis Martínez Mont, Professor of Experimental Psychology in Secondary Schools of State.

His Majesty the King of Hejaz, Nejd and Dependencies:

Cheik Hafiz Wahba, Envoy Extraordinary and Minister Plenipotentiary to His Britannic Majesty.

His Majesty the King of Italy:

M. Stefano CAVAZZONI, Senator, Former Minister of Labour,

His Majesty the Emperor of Japan;

M. Setsuzo Sawada, Minister Plenipotentiary, Director of the Japanese Bureau accredited to the League of Nations;

M. Shigeo Ohdachi, Secretary at the Ministry for Home Affairs, Head of the Administrative Section.

The President of the Republic of Liberia.

「ドクトル、アントアンス、ソッティール」國際聯盟ニ派遣ノ常任代表委員、特命全權公使

外務方課長「ジャオザス、サカラウスカス」外務大臣「ドクトル、ドヴァス、ザウニウス」「リスアニア」共和國大統領

「ルクセンプルグ」國大公殿下

在「ジュネーグ」簡単「シァググ、グェグメーグ」

「メキシュ」合衆國大統領

マルティネス・ディアルヴァ」國際聯盟ニ派遣ノ常任傍聴委員「サルヴァドル、

「モナコ」國公殿下

在「ジュネーグ」熱領事「コンラド、エー、アンチ」

「パナマ」共和國大統領

在「ジュネーツ」総領事「ドクトル、エルネスト、

「パラグァイ」共和國大統領

國際聯盟ニ版造ノ常任代表委員、佛蘭西共和國

Dr. Antoine Sottile, Envoy Extraordinary and Minister Plenipotentiary, Permanent Delegate accredited to the League of Nations.

The President of the Republic of Lithuania:

Dr. Dovas Zaunius, Minister for Foreign Affairs.

M. Juozas Sakalauskas, Head of Section at the Ministry for Foreign Affairs.

Her Royal Highness the Grand-Duchess of Luxemburg:

M. Charles VERMAIRE, Consul at Geneva.

The President of the United States of Mexico:

M. Salvador Martínez de Alva, Permanent Observer accredited to the League of Nations.

His Serenc Highness the Prince of Monaco:

M. Conrad E. Hentsch, Consul-General at Geneva.

The President of the Republic of Panama:

Dr. Ernesto Hoffmann, Consul-General at Geneva.

The President of the Republic of Paraguay:

Dr. Ramón V. CABALLERO DE BEDOYA, Envoy

カバリェロ、ディベドャ」 駐箚特命全権公使「ドクトル、ラモン、ヴェー、

和蘭國皇帝陛下

ファン、ヴェットゥムー 関際阿片問題 二開スル 政府顧問「ヴェー"()、

「イグット」医型作器下

特命全権公使「エー、セパーボディ」國際聯盟ニ派遣ノ常任代表委員、瑞西聯邦駐箚

前大臣「ヴィトルド、キデュ」「ボーランド」共和國大統領

「ボルトガル」共和國大統領

、ル、アウグスト、デ、ヴァスコンセロスー「ボルトガル」國國際聯盟局長、全権公使「ドクト

ス、ディアンドラーデ」書記官「ドクトル、アレシァンドロ、フェルラ関際聯盟「ボルトガル」國事務局長、公使館一等

「ガートニア」國皇帝陛下

Extraordinary and Minister Plenipotentiary to the President of the French Republic, Permanent Delegate accredited to the League of Nations.

Her Majesty the Queen of the Netherlands:

M. W. G. VAN WETTEM, Government Adviser for International Opium Questions.

His Imperial Majesty the Shah of Persia:

M. A. Sepaibody, Envoy Extraordinary and Minister Plenipotentiary to the Swiss Federal Council, Permanent Delegate accredited to the League of Nations.

The President of the Polish Republic:

M. Witold Cποτέκο, Former Minister.

The President of the Portuguese Republic:

Dr. Augusto de Vasconcellos, Minister Plenipotentiary, General Director of the Portuguese Secretariat of the League of Nations;

Dr. Alexandro Ferraz de Andrade, First Secretary of Legation, Chief of the Portuguese Office accredited to the League of Nations.

His Majesty the King of Roumania:

ティン、アントニアーデ」國際聯盟ニ 派 造 ノ 特命全権公使「コンスタン

「サン、マリノ」共和國攝政官

辯護士、教授「チー、エー、フェルリー

遇羅國皇帝生下

権公使「ダムラス」戦下國際聯盟ニ派遣ノ常任代表者、英國駐箚特命全

路典屬學術籍下

トマン」 瑞西聯邦駐箚特命全権公使 「コーパイー、ヴェス

路西縣邦政府

ジェール」 聯邦政務省外務部長、全権公使「ボール/ディニ

エール」
聯邦公衆衛生部長「ドクトル、アンリー、カリ

「チェッコスロヴァキア」共和國大統領

國際聯盟三派遣ノ常任代表委員、瑞西聯邦駐箚

M. Constantin Antoniade, Envoy Extraordinary and Minister Plenipotentiary accredited to the League of Nations.

I Capitani Reggenti of the Republic of San Marino:

Professor C. E. FERRI, Advocate.

His Majesty the King of Siam:

His Serene Highness Prince Damras, Envoy Extraordinary and Minister Plenipotentiary to His Britannic Majesty, Permanent Representative accredited to the League of Nations.

His Majesty the King of Sweden:

M. K. I. Westman, Envoy Extraordinary and Minister Plenipotentiary to the Swiss Federal Council.

The Swiss Federal Council:

M. Paul DINICHERT, Minister Plenipotentiary, Chief of the Division for Foreign Affairs of the Federal Political Department;

Dr. Henri CARRIERE, Director of the Federal Service of Public Health.

The President of the Czechoslovak Republic:

M. Zdeněk Fierlinger, Envoy Extraordinary

ル」 特命全権公使「ズデニェック、フィエルリング

「ウルグァイ」共和國大統領

ド、デ、カストロ」 瑞西聯邦 駐街解析 生衛代金を「ドクトル、アルフレ

「ヴェネズエラ」合衆國大統領

クトル、エレ、ヘー、チァシン・イトリアゴー「カラカス」際學院會員、在「ベルス」代理公使「ド

安當ナルヲ認メタル後左ノ如ク協定セリ右各全権委員、五二共ノ全権委任张ラ示シ之が良好

第一章定義

按 1 些

通ジ適用セラルベシ別段ノ明示アル場合ヲ除クノ外左記定義へ本條約ヲ

グー接利

約ヶ謂フ 日「ジュネーツ」」、於テ緊名セラレタル 國際阿片條一、ジュネーツ」は終約トハ 千九百二十五年二月十九 and Minister Plenipotentiary to the Swiss Federal Council, Permanent Delegate accredited to the League of Nations.

The President of the Republic of Uruguay:

Dr. Alfredo DE CASTRO, Envoy Extraordinary and Minister Plenipotentiary to the Swiss Federal Council.

The President of the United States of Venezuela:

Dr. L. G. Chacin-Itrhago, Chargé d'Affaires at Berne, Member of the Medical Academy of Caracas.

Who, having communicated to one another their full powers, found in good and due form, have agreed as follows:

CHAPTER I.—DEFINITIONS.

ARTICLE 1.

Except where otherwise expressly indicated, the following definitions shall apply throughout this Convention:

The term "Geneva Convention" shall denote the International Opium Convention signed at Geneva on February 19th, 1925.

問いズ左記薬品ヲ謂フ栗 品 二 薬品トい字製ノモノナルト精製ノモノナルトラ

第1種

亞麵人

- ル「モルとネ」ラ含有スル製劑ラ含ム) 片ョリ直接作ラレ且二〇「パーセント」ヲ超ユ 一「モルとネ」及其ノ鹽類(生阿片又、藥用阿
- ステル」並二其ノ鹽類「デアセチルモルヒネ」及他ノ「モルヒネエ
- ゴニンエステル」及其ノ鹽類と」う含有スル製剤う含ム) 並ニ一切ノ「エク作ラレ且○・」「バーセント」ヲ超エル「コカイ国・「ニカイン」 及其ノ鹽類(「コカ」薬=リ直接
- 類ナリ)、「アセチルデヒドロコディノン」即り」と名ノ下ニ 登録セラレタル物質が其ノ躍類ナリ)、「デヒドロモルヒノン」(「デラウデット」く名ノ下ニ 登録セラレタル 物質が其ノ躍類ナリ)、「デヒドロコディノン」(「デコデッル」く名ノ下ニ登録セラレタル 物質が其ノ躍

2. The term "the drugs" shall denote the following drugs whether partly manufactured or completely refined:

Group I.

Sub-Group (a):

- (i) Morphine and its salts, including preparations made directly from raw or medicinal opium and containing more than 20 per cent of morphine;
- (ii) Diacetylmorphine and the other esters of morphine and their salts;
- (iii) Cocaine and its salts; including preparations made direct from the coca leaf and containing more than 0.1 per cent of cocaine, all the esters of ecgonine and their salts;
- (iv) Dihydrohydrooxycodeinone (of which the substance registered under the name of eucodal is a salt); dihydrocodeinone (of which the substance registered under the name of dicodide is a salt); dihydromorphinone (of which the substance registered under the name of dilaudide is a salt), acetyldihydrocodeinone or acetyldemethylodihydrothebaine (of which the substance registered under the

チーアセチルディチロデロティイン」(「ア セデコン」ノ名ノ下ニ 登録セラレタル 物質 パ **実ノ堕類ナリ)、「デヒドロモルヒネ」(「パラモ** ルファン」ノ名ノ下二登録セラレタル物質の其 ノ踵類ナリ)、共ノ「エステル」、右物質ノ何レ カノ鹽類及其ノ「エステル」ノ鹽類、「モ州ヒ ネ・エス・オキシード」(登録名「ゼノモルヒネ」) 立二「モルとネ・エヌ・オキシード」 誘導體及他 ノ五價窒素「モルヒネ」誘導體

田屋口

「エクゴニン」、「テパイン」及共ノ鹽類拉ニ「ベ ンジルモルヒネ」、他ノ「モルヒネエーテル」 及其ノ鹽類但シ「メチルモルヒネ」(「コデイ ン」)、「エチルモルヒネ」及其ノ鹽類ヲ除ク

「メチグモグロネ」(「コディン」)、「エチグモグロ ネ」及 共ノ 鹽 類

本號二記載セラルル物質い合成ノ方法二佐り製産 セラルルトキト雖モ藥品ト育做サル

第一類及第二類トハ夫々本號ノ第一類及第二類ラ

name of acedicone is a salt); dihydromorphine (of which the substance registered under the name of paramorfan is a salt), their esters and the salts of any of these substances and of their esters, morphine-N-oxide (registered trade name genomorphine), also the morphine-N-oxide derivatives, and the other pentavalent nitrogen morphine derivatives.

Sub-Group (b):

Ecgonine, thebaine and their salts, benzylmorphine and the other ethers of morphine and their salts, except methylmorphine (codeine), ethylmorphine and their salts.

Group II. .

Methylmorphine (codeine), ethylmorphine and their salts.

The substances mentioned in this paragraph shall be considered as drugs even if produced by a synthetic process.

The terms "Group I" and "Group II" shall

馬フ

生 岡 片 二二 生阿片トハ罌巣(「パパヴェ、フムニフェラム、エ ル」) 賞ヨリ 得タル 液汁ノ自然ニ凝結シタルモノ ニシテ「モルヒネ」合行量ノ如何ヲ問ハズ耶ニ包裝 及輸送ニ必要ナル程度ノ加工ラ為シタルモノラ間

聚田区土

薬用阿片ト、粉状、粒状义、他ノ 形状ノモノタル ト中性物ヲ混ズルモノタルトヲ問ハズ内國藥局方 ノ定ムル所ニ從と路藥用ニ適應セシムルニ必要ナ ル加工ラ為シタル生阿片ラ腊フ

「モルヒネ」トハ阿片ノ主要ナル「アルカロイド」ニ シテ C17H19O5パノ化學式ラ行スルモノラ謂フ 「米トヤルシャシンド」~ 、 C21H23O5N(C17H17(C2 H3O)2O3X/ 人化原式ラ行スル「デアセチルモルと ネ」(「デアモルとネ」、「ヘロイン」) ラ腊フ

「ロセ」類

「ヨカ」葉トハ古加樹科ニ協スル「エリトロキシロ ン、スカ、ラマグク」、「エリトロキッロン、ノグナ・グ ラナテンス(モリス)、ヒエロニムス」及其ノ緑種ノ 薬竝ニ右属ノ他ノ種ノ薬ニシテ之ヨリ直接又ハ化 respectively denote Groups I and II of this paragraph.

3. "Raw opium" means the spontaneously congulated juice obtained from the capsules of the Papaver somniferum L., which has only been submitted to the necessary manipulations for packing and transport, whatever its content of morphine.

"Medical opium " means raw opium which has undergone the processes necessary to adapt it for medicinal use in accordance with the requirements of the national pharmacopæia, whether in powder form or granulated or otherwise or mixed with neutral materials.

"Morphine" means the principal alkaloid of opium having the chemical formula C17H19O3N.

"Diacetylmorphine" means diacetylmorphine (diamorphine, heroin) having the formula $C_{21}H_{23}$ O5N (C17H17(C2H3O)2O3N).

'Coca leaf" means the leaf of the Erythroxylon Coca Lamarck and the Erythroxylon novogranatense (Morris) Hieronymus and their varieties, belonging to the family of Erythroxylaceæ and the leaf of other species of this genus from which it may be found possible to extract cocaine,

ノ五價窒素「モルヒネ」誘導體及他立「モルヒネ・エス・オキシード」誘導體及他ネ・エス・オキシード」(登録名「ゼノモルヒネ」)カノ鹽類及其ノ「エステル」ノ鹽類、「モルヒノ鹽類ナリ)、其ノ「エステル」、右物質ノ何レルファン」ノ名ノ下ニ登録セラレタル物質、其よの題類ナリ)、「デヒドロモルヒネ」(「バラモセデコン」ノ名ノ下ニ登録セラレタル物質、其

田類日

ン」)、「エチルモルヒネ」及其ノ鹽類ヲ除ク及其ノ鹽類但シ「メチルモルヒネ」(「コデインジルモルヒネ」、他ノ「モルヒネエーテル」「エクゴニン」、「テバイン」及其ノ鹽類立ニバ

班11年

ネ」及状ノ騒灯「メチルモルとネ」(「コデイン」)、「エチルモルと

セラルルトキト雖モ藥品ト看做サル本跳ニ記載セラルル物質い合成ノ方法ニ佐り製産

第一類及第二類トハ夫々本號ノ第一類及第二類ラ

name of acedicone is a salt); dihydromorphine (of which the substance registered under the name of paramorfan is a salt), their esters and the salts of any of these substances and of their esters, morphine-N-oxide (registered trade name genomorphine), also the morphine-N-oxide derivatives, and the other pentavalent nitrogen morphine derivatives.

Sub-Group (b):

Ecgonine, thebaine and their salts, benzylmorphine and the other ethers of morphine and their salts, except methylmorphine (codeine), ethylmorphine and their salts.

Group II. .

Methylmorphine (codeine), ethylmorphine and their salts.

The substances mentioned in this paragraph shall be considered as drugs even if produced by a synthetic process.

The terms "Group I" and "Group II" shall

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田阿片

ス輪送ニ必要ナル程度ノ加工ヲ為シタルモノヲ謂ニシテ「モルヒネ」含有量ノ如何ヲ問ハズ罪ニ包裝ル」) 賞ヨリ 得タル 液汁ノ自然ニ凝結シタルモノ三 生阿片トハ器架(「ババヴェ`ソムニフェラム、エ

聚田厚土

ル加工ヲ爲シタル生阿片ヲ謂フノ定ムル所ニ從と醫藥用ニ適應セシムルニ必要ナ;中性物ヲ混ズルモノタルトヲ問ニズ内國藥局方藥用阿片トハ粉狀、粒狀父ハ他ノ形状ノモノタル

ネ」(「デアモルヒネ」、「ヘロイン」) ヲ謂フH³O)²O³Z) ~化原式ヲ行スル「デアセチルモルヒ「デアセチルモルとよ」トハ C²1H²3O²Z(C¹7H¹7(C²シテ C¹2H¹9O³Z ~化學式ヲ行スルモノヲ謂フ「モルヒネ」トハ阿片ノ主要ナル「アルカロイド」ニ

「ロセ」数

薬竝ニ右属ノ他ノ種ノ薬ニシラ之ヨリ直接父ハ化ラナテンス(モリス)、ヒエロニムス」及其ノ髪種ノン、ヨカ、ラマルク」、「エリトロキシロン、ノヴォグ「コカ」葉トハ古加樹科ニ協スル「エリトロキシロオ」(「チアモルヒネ」、「(ロイン」)ヲ謂フ

respectively denote Groups I and II of this paragraph.

3. "Raw opium" means the spontaneously coagulated juice obtained from the capsules of the Papaver somniferum L., which has only been submitted to the necessary manipulations for packing and transport, whatever its content of morphine.

"Medical opium" means raw opium which has undergone the processes necessary to adapt it for medicinal use in accordance with the requirements of the national pharmacopæia, whether in powder form or granulated or otherwise or mixed with neutral materials.

"Morphine" means the principal alkaloid of opium having the chemical formula C₁₇H₁₉O₃N.

"Diacetylmorphine" means diacetylmorphine (diamorphine, heroin) having the formula $C_{21}H_{23}$. O_5N ($C_{17}H_{17}(C_2H_3O)_2O_3N$).

"Coca leaf" means the leaf of the Erythroxylon Coca Lamarck and the Erythroxylon novogranatense (Morris) Hieronymus and their varieties, belonging to the family of Erythroxylaceæ and the leaf of other species of this genus from which it may be found possible to extract cocaine,

堕的方法 | 依り「コカイン」ヲ抽出スルコトヲ得べ しょかイソート シュイチ・ベングイグ、フータ・ナー 「ロセナン」 クゴョン」(110パーセント」「クロロギブム」溶液 ニ於ケル比旋光度 ([a] D20°) 左旋一六・四度) ニシ テ CivH21O4N ~化學式?有スルモノラ謂ブ 「エクゴニン」ト ハ「フーヴォ・エクゴニン」(五「パー セント」水溶液ニ於ケル 比薩光度 ([a] D20°) 左旋 四五・六度)ニシテ CoH15O3N.H2O~化學式ラ有ス ルモノ及工業上「レーザナ・エクゴニン」ノ再製二供 シ得べキ 一切ノ「レーヴォエクゴニン」誘導體ラ語 左ノ藥品の左記化學式ニ依り之う定ム 「デヒドロオキシコデイ ${\textstyle \nwarrow \lambda \rfloor \ \cdots \cdots C_{18} H_{21} O_4 N}$ 「米 1 1 1 1 1 1 1 イ / 入」 ······ C18 H21 O3 N 「アセチガデンドロコデ イノン」又ハ「アセチル アメチロチャットロティ

either directly or by chemical transformation

"Cocaine" means methyl-benzoyl laevo-ecgonine ([a] D $20^\circ = -16^\circ 4$) in 20 per cent solution of chloroform or which the formula is $C_{17}H_{21}O_4N$.

"Ecgonine" means lagevo-ecgonine ([a] D 20° = -45°6 in 5 per cent solution of water), of which the formula is C₉H₁₅O₃N.H₂O, and all the derivatives of laevo-ecgonine which might serve industrially for its recovery.

The following drugs are defined by their chemical formulæ as set out below: Dihydrohydrooxycodeinone... $C_{18}H_{21}O_4N$

	$\begin{array}{cccccccccccccccccccccccccccccccccccc$
	「米 n * p # × n *」 · · · · · · · C ₁₇ H ₂₁ O ₃ N
	「モルとネ・エス・オキシー
	½
	「水 ~ ~)」 · · · · · · · · · · · · · · · · · ·
	「メチガホガカ头」(「ログ
	$\begin{array}{c} \leftarrow \ \searrow \ \rfloor) \cdots $
8	$ \begin{array}{c} \Gamma \not H \not + \not \geq \psi \not \geq \omega \not \leftarrow \rfloor \cdots \cdots C_{19} H_{23} O_3 N(C_{17} H_{18} \\ (C_2 H_5 O) O_2 N) \end{array} $
	「ツスジミルミル・」 $C_{24}H_{25}O_{3}N(C_{17}H_{18}$ ($C_{7}H_{7}O)O_{2}N$)
彩 朋	四 製造ト・、精製ノ何レノ過程ラモ包含ス
* *	轉換トハ化學的方法ニ依ル藥品ノ變形ヲ謂フ但シ
	「アルカロイド」ラ北ノ騒刺ニ 秘形スル コトラ除
	一ノ藥品が他ノ薬品ニ轉換セラルル場合其ノ操作
	(前者二對スル開係二於テ、轉換ニシテ後者上對一分勇品力化 季品 一十
	スル開係二次テ、製造ナリト行俄サル・前子一造ラル開係になった。

	$(C_2H_3O)O_3N)$
Dihydromorphine	C ₁₇ H ₂₁ O ₃ N
Morphine-N-Oxide	
Thebaine	C ₁₉ H ₂₁ O ₃ N
Methylmorphine (codein	$\stackrel{\text{(e)}}{\text{(cH_{3}O)O_{2}N}} \stackrel{\text{(cH_{18}H_{21}O_{3}N(C_{17}H_{18}))}}{\text{(cH_{3}O)O_{2}N)}}$
Ethylmorphine	$C_{19}H_{23}O_3N(C_{17}H_{18} - (C_2H_5O)O_2N)$
Benzylmorphine	$O_{24}H_{25}O_3N(C_{17}H_{18})$ $(C_7H_7O)O_2N)$
any process of refining	'shall denote the trans- chemical process, with

the exception of the transformation of alkaloids

When one of the drugs is converted into another of the drugs this operation shall be considered as conversion in relation to the firstmentioned drug and as manufacture in relation to

into their salts.

the other

D! :

見償ヲモ包含スルルと視す謂と且文意ニ別段ノ要求ナキ限リ循足見積トハ本條約第二條方至第五條ニ從と提用セラ

全国内证明

スル信仰品で謂う謂う事倫任印品トハ在記ノ為ニ要何レカノ樂品ニ闢と準備任印品トハ在記ノ為ニ要

- 内部官役(在庫品)保存セラルル國文・領域・正常ナル
- 二 右ノ國文(領域三於ケツ種換

111 3建田

セラルル作庫品ヲ謂フ啓及例外的事情ニ順ズル為政府ノ取縮ノ下ニ保存何レカノ薬品ニ關シ政府作庫品トハ政府ノ使用ノ

攀田

スルモノト行後サル文章ニ別段ノ要求ナキ限り輸出、再輸出プモ包含

第二位品質

孫 11 秦

可能人思

各領域ニ腸シ 本族約 第五條ノ 規定ニ依ル 見蹟ラー 各締約國ハ各藥品ニ件本條約ノ適用アル自國ノ

The term "estimates" shall denote estimates furnished in accordance with Articles 2 to 5 of this Convention and, unless the context otherwise requires, shall include supplementary estimates.

The term "reserve stocks" in relation to any of the drugs shall denote the stocks required

- (i) For the normal domestic consumption of the country or territory in which they are maintained,
- (ii) For conversion in that country or territory, and
- (iii) For export.

The term "Government stocks" in relation to any of the drugs shall denote stocks kept under Government control for the use of the Government and to meet exceptional circumstances.

Except where the context otherwise requires, the term "export shall be deemed to include re-export

CHAPTER II.—ESTIMATES.

ARTICLE 2.

 Each High Contracting Party shall furnish annually, for each of the drugs in respect of each

設中央委員會ニ毎年提出スペシ「ジュネーツ」條約第六章ニ依り 改置セラレタル常

- 1祝グラルル監督機關二依り提出セラルベシ提出セザル場合二ハ見積、能フ限り第五條第大號三開シ見積フ第五條第四號二祝グラルル期日迄二二一締約國ガ木條約ノ適用アル自國ノ何レカノ領域
- 積ヲ作成スペシ 提出セラレザルトキ・監督機關・育ラ能フ限リ兄ヲ要求スペシ右ノ何レカノ國父・領域ニ付見積ガニ付見積ガ本條約ノ規定ニ從と作成セラルルコト三 常設中央委員合・本條約ノ適用ナキ國父・領域

统 三族

是一口流

ルコトラ得見取りとうかな事情と説明と共ニ提出スと初域ニ闘シ談年ニ付テノ談領域ニ闘スル補足総約國人の要ナルトキハ何レノ年ニ於テモ自國ノ何

of his territories to which this Convention applies, to the Permanent Central Board, constituted under Chapter VI of the Geneva Convention, estimates in accordance with the provisions of Article 5 of this Convention.

- 2. In the event of any High Contracting Party failing to furnish, by the date specified in paragraph 4 of Article 5, an estimate in respect of any of his territories to which this Convention applies, an estimate will, so far as possible, be furnished by the Supervisory Body specified in paragraph 6 of Article 5.
- 3. The Pemanent Central Board shall request estimates for countries or territories to which this Convention does not apply to be made in accordance with the provisions of this Convention. If for any such country estimates are not furnished, the Supervisory Body shall itself, as far as possible, make the estimate.

ARTICLE 3. .

Any High Contracting Party may, if necessary, in any year furnish in respect of any of his territories supplementary estimates for that territory for that year with an explanation of the circumstances which decessitate such supplementary estimates.

张 回 继

見隋〉还衛

抜クベシリ事ラ右ノ國文・領域ノ際採用及學術用ノ霊要ニ**國文**・領域ノ内部消費ニ必要ナル藥品ニ關スル限一前諸株ニ従ヒ提出セラルル各見稿・其ノ關スル

で変なが出

保有スルコトラは11一緒約國ハ準備在庫品ノ外政府在庫品ヲ設定シ且

第 五 條

日籍人比式

メラルル 非韓盟國ニ道知 モラルル様式 二健フ ベニ依り國際聯盟ノ一切ノ聯盟國及第二十七條三拐「當眾中央委員合三依り隨時定メラレ且同委員會」 本條約第二條乃至第四條42規定セラルル各是額

りにはよく

- 、ズ各薬品ニ腸シ左記ラ示スペシロイド」又、鹽額ノ製剤ノ形態ノモノタルトラ問ルカロイド」又、鹽額ノ形態ノモノタルト「アルカレイド」又、鹽額ノ形態ノモノタルト「アルカフ、各見積ニ、各國又、各領域ニ付及各年ニ付「ア
 - 降塚用及學術用ノ為共レ自體トシテノ使用ニ

ARTICLE 4.

- Every estimate furnished in accordance with the preceding Articles, so far as it relates to any of the drugs required for domestic consumption in the country or territory in respect of which it is made, shall be based solely on the medical and scientific requirements of that country or territory.
- The High Contracting Parties may, in addition to reserve stocks, create and maintain Government stocks.

ARTICLE 5.

- 1. Each estimate provided for in Articles 2 to 4 of this Convention shall be in the form from time to time prescribed by the Permanent Central Board and communicated by the Board to all the Members of the League of Nations and to the non-member States mentioned in Article 27.
- Every estimate shall show for each country or territory for each year in respect of each of the drugs whether in the form of alkaloid or salts or of preparations of the alkaloids or salts;
 - (a) The quantity necessary for use as

ヲ要セザル製剤ノ製造ニ必要ナル数量ヲ含ム)ノ偽ノモノタルトヲ問ハズ輸出スルニ輸出許可必要ナル数量(内部消費ノ為ノモノタルト輸出

- ルトラ問へズ轉換用こ必要ナル数量の 内部消費ノ為ノモノタルト輸出ノ為ノモノタルト輸出ノ為ノモノタ
- 3 保行セント欲スが準備在庫品ノ数量
- 有二必要ナル数量 第四條二規定セラルル政府在庫品ノ設定及保

開内ニ近付セルニ非ザル限り考慮セラレザルベシ約岡方常設中央委員合ニ必要ナル見領ヲ適當ノ期入ヨヲ成ルベシ但シ此等ノ加算又、投除、關係緒平準ヲ超過スルコトアルベキ數量ヲ控除シタルモベキ數量ヲ加算シ又、右合計ヲ化ニ必要ナルコトアルニる要ノルニを建ニ達セシムルニ必要ナルコトアルニ協グラルル數量ノ合計ニ準備在庫品及政府在庫1級又、各領或「付テノ見積ノ總量、本號(及回

such for medical and scientific needs, including the quantity required for the manufacture of preparations for the export of which export authorisations are not required, whether such preparations are intended for domestic consumption or for export;

 (b) The quantity necessary for the purpose of conversion, whether for domestic consumption or for export;

(c) The amount of the reserve stocks which it is desired to maintain;

(d) The quantity required for the establishment and maintenance of any Government stocks as provided for in Article 4.

The total of the estimates for each country or territory shall consist of the sum of the amounts specified under (a) and (b) of this paragraph with the addition of any amounts which may be necessary to bring the reserve stocks and the Government stocks up to the desired level, or after deduction of any amounts by which those stocks may exceed that level. These additions or deductions shall, however, not be taken into account except in so far as the High Contracting Parties concerned shall have forwarded in due course the necessary estimates to the Permanent Central Board.

見職へ発明

ナルコトアリ得ルモノトス
、他ノ藥品ノ場合ニ於ケルヨリ大ナル餘裕ノ必要セラルルコトアルベキ何レカノ藥品ノ場合ニ於テ星ヲ指示スルヲ要ス第二類ニ包含セラレ又、包含ラレタルトキハ見積ニ、斯ク包含セラレタル餘裕有り得べキ變動ニ對スル餘裕ヲ包含スル樣計算セレタル方法ノ説明書ヲ添附スベシ右數量が需要ノ」各見荷ニハ之ニ記載セラルル諸數量が計算セラ

関係~後田

常設中央委員會ニ到達スルコトラ要ス四、各見荷、其ノ關スル年ノ前年ノ八月一日以前ニ

関于日本

付セラルベシ玉 補足見積い其ノ完了後直ニ常設中央委員會ニ送

員會トノ密接ナル協力ラ確保スペシ珍總長「常設中央委務長」依り設ケラルベク事務總長「常設中央委ノ権利ヲ有スペシ監督機關ノ事務局「國際聯盟事衛生國際事務局「各右監督機關ノ一員ヲ任命スル合、常設中央委員會、國際聯盟保健委員會立一公案」「阿片及他」危險藥品ノ取引二關スル 諮問委員大 見荷い監督機關」依り檢査セラルベシ國際聯盟

- 3. Every estimate shall be accompanied by a statement explaining the method by which the several amounts shown in it have been calculated. If these amounts are calculated so as to include a margin allowing for possible fluctuations in demand, the estimates must indicate the extent of the margin so included. It is understood that in the case of any of the drugs which are or may be included in Group II, a wider margin may be necessary than in the case of the other drugs.
- Every estimate shall reach the Permanent Central Board not later than August 1st in the year preceding that in respect of which the estimate is made.
- Supplementary estimates shall be sent to the Permanent Central Board immediately on their completion.
- 6. The estimates will be examined by a Supervisory Body. The Advisory Committee on the Traffic in Opium and other Dangerous Drugs of the League of Nations, the Permanent Central Board, the Health Committee of the League of Nations and the Office international d'Hygiène publique shall each have the right to appoint one member of this Body. The Secretariat of the Supervisory Body shall be provided by the Sec-

和如你里

没明ニテ足ルモノトス 及用ニチアルベキ何レカノ藥品」場合ニ於テ、散略スルコトヲ得第二額ニ包含セラル又、包含セラル又、詳細ニ基キ關係政府ノ同意ヲ以テ見預ヲ修正細ヲ災ニ要求スルヲ得ベク且斯クシテ得タル情報ニ共ノ必要ナリト認ムルコトアルベキ情報又、詳古ニシメ又、之ニ記載セラルル非項ヲ武明スル為提用セラレタル國又、領域ニ關ス、法見荷ヲ完全監督機關、政府ノ雷要ニ關スルモノヲ除キ見積ノ

屋 上

京セラレタル説明ノ要領及監督機開ガ右見積若いりト認とが限り前記第大號ニ従と與ヘラレダ、要及ハ各領域ニ付ラノ 見積ノ表、監督機關ガ必要ナ聯盟國及第二十七條ニ協グラルル非聯盟國ニ各國月一日以前ニ非務總長ヲ総由シ國際聯盟ノ一切ノニ條ニ規定セラルル決定ノ後監督機關、每年十一次各國文、各領域ニ付ラノ見積ノ右機關ニ佐ル第二級にセラレッ協在ノ後及見積ノ結出セラレザ社 提出セラレタル見積ノ監督機關ニ佐ル前記第六

retary-General of the League of Nations, who will ensure close collaboration with the Permanent Central Board.

The Supervisory Body may require any further information or details, except as regards requirements for Government purposes, which it may consider necessary, in respect of any country or territory on behalf of which an estimate has been furnished in order to make the estimate complete or to explain any statement made therein, and may, with the consent of the Government concerned, amend any estimate in accordance with any information or details so obtained. It is understood that in the case of any of the drugs which are or may be included in Group II a summary statement shall be sufficient.

7. After examination by the Supervisory Body as provided in paragraph 6 above of the estimates furnished, and after the determination by that Body as provided in Article 2 of the estimates for each country or territory on behalf of which no estimates have been furnished, the Supervisory Body shall forward, not later than November 1st in each year, through the intermediary of the Secretary-General, to all the Members of the League of Nations and nonmember States referred to in Article 27, a statement containing the estimates for each country or

コトアルベキ意見ヲ恣付スベシ説明又ハ説明ノ要求ニ關シ其ノ表明セント欲スル

と監督機關ニ佐り延備ナク處理セラルベシ足額、前記第六號及第七號ニ祝デリがフルル手額ニ従入 年中ニ常設中央委員會ニ送付セラレタル各補足

第 三 章 製造網膜

第六條

Manual

- ラルルコトナカルベシノ薬品ノ飲益い左記數益ノ合計ヲ超過シテ製造セー何レノ闽文、領域ニ於テモ一年間ニ於テ何レカ
- 一輪出許可ヲ要セザル製剤ノ製造」必要ナル数タルト輸出ノ為ノモノタルトヲ問ハズ輸出スルテノ使用」必要ナル数量(内部消費ノ為ノモノ無関内」於子醫及用及學術用ノ為我レ自體トッイ年二付テノ右ノ國及ハ領域ニ關スル見積ノ

territory and, so far as the Supervisory Body may consider necessary, an account of any explanations given or required in accordance with paragraph 6 above, and any observations which the Supervisory Body may desire to make in respect of any such estimate or explanation, or request for an explanation.

8. Every supplementary estimate sent to the Permanent Central Board in the course of the year shall be dealt with without delay by the Supervisory Body in accordance with the procedure specified in paragraphs 6 and 7 above.

CHAPTER III .- LIMITATION OF MANUFACTURE.

ARTICLE 6.

- 1. There shall not be manufactured in any country or territory in any one year a quantity of any of the drugs greater than the total of the following quantities:
 - (a) The quantity required within the limits of the estimates for that country or territory for that year for use as such for its medical and scientific needs including the quantity required for the manufacture of preparations for the export of which export

展す合(4)

- 為ノモノタルトヲ問ハズ轉換ニ必要ナル数量範圍内ニ於テ内部消費ノ為ノモノタルト輸出ノロ「右年ニ付テノ右ノ國又、領域ニ關スル見積ノ
- キ戦量 スル為右ノ國父、領域ガ必要トスルコトアルベス 本條約ノ規定ニ依ル輸出ノ註文ヲ年内ニ城行
- コトアルベキ数点 平準二維持スル為右ノ岡又ハ領域ガ必要トスルー 準備作用品ヲ右年ニ付テノ見積ニ掲グラルル
- 平準ニ維持スル局必要トスルコトアルベキ數量大 政府作庫品ラ右年ニ付テノ見積ニ掲グラルル

スル音句の問うな

ルベキモノトス締約國八常設中央委員會二自國ノ過量、墾年中二製造セラルベキ數量ヨリ控除セラルコトヲ締約國ガ年末二於テ發見スルトキハ右超作第一項二依ル羟除ヲ為シテ得タル數益ヲ超過ス1一製造セラレタル數益ガ前記數量ノ合計ヨリ第七

authorisations are not required, whether such preparations are intended for domestic consumption or for export;

- (b) The quantity required within the limits of the estimates for that country or territory for that year for conversion, whether for domestic consumption or for export;
- (c) Such quantity as may be required by that country or territory for the execution during the year of orders for export in accordance with the provisions of this Convention.
- (d) The quantity if any, required by that country or territory for the purpose of maintaining the reserve stocks at the level specified in the estimates for that year;
- (e) The quantity, if any, required for the purpose of maintaining the Government stocks at the level specified in the estimates for that year.
- 2. It is understood that, if at the end of any year, any High Contracting Party finds that the amount manufactured exceeds the total of the amounts specified above, less any deductions made under Article 7, paragraph 1, such excess shall be deducted from the amount to be manufactured during the following year. In forwarding their

年次統計ラ送付スルニ常り右超過ノ理由ラ示スペ

第七條

キ敷削 応セラル(機関マヨリな 各種の東田製造

長ヲ控除スペシニャヲ許サルル各藥品ノ總量ヨリ左記数第六後ニ佐リ何レカノ國及ハ領域ニ於テ一年間ニ製

- 再輸出量ヲ控除シテ得タル數量 | 返還セラレタルモノヲ含ム輸入藥品ノ數量ョリ
- ラルベシ末ニ於テ殘存スル超過數量、翌年ノ見積ヨリ控除と當該年中ニ前記控除ノ何レカヲ為シ得ザルトキハ年消費ノ為父ハ轉換ノ為ニ利用セラルルモノノ數是□□ 押牧セラレタル薬品ニシテ共レ自體トシテ内部

第 八 條

整造ライ制の 日本には映画 イラスペート 全ルベ人目従 数ル製を向し

ルル何レカノ薬品ノ敷量、可能ナルトキ、右見積ノノ為該國义、該領域ニ於テ輸入セラレ又、製造セラ何レカノ國文、領域ニ件テノ見積ニ従と轉換ノ目的何レカノ國文、領域ニ件テノ見積ニ従と轉換ノ目的

annual statistics to the Permanent Central Board, the High Contracting Parties shall give the reasons for any such excess.

ARTICLE 7.

There shall be deducted from the total quantity of each drug permitted under Article 6 to be manufactured in any country or territory during any one year:

- (i) Any amounts of that drug imported including any returned deliveries of the drug, less quantities re-exported.
- (ii) Any amounts of the drug seized and utilised as such for domestic consumption or for conversion.

If it should be impossible to make any of the above deductions during the course of the current year, any amounts remaining in excess at the end of the year shall be deducted from the estimates for the following year.

ARTICLE 8.

The full amount of any of the drugs imported into or manufactured in any country or territory for the purpose of conversion in accordance with the estimates for that country or territory shall,

動を田

返用セラルル期間内 1 右目的ノ為全部利用セラルベ

数は部分へ

該ヨリ珍除セラルベシスル部分の選年ニ付テノ右ノ岡父の領域ニ闘スル見可能ナル場合ニハ年末ニ於テ利用セラレズシテ選作尤モ右期間内ニ右目的ノ為全数基ヲ利用スルコト不

第九條

在議品人超

り控除セラルベシ合ニ腫ジ通常輸入セラレス、製造セラルベキ數量ョハ該超過量、常該年中ニ於ラ本條約ノ規定ニ依り場ト欲スル該藥品ノ準備任庫品ン數量ヲ超過スルトキ庫品ガ該國又、該領域ニ付テノ見額ニ依り保有セン又、領域ニ於ケル何レカノ藥品ノ共ノ常時ノ現存任本條約ノ一切7規定が宜施セラレタル際何レカノ國

が數量、該年中場合ニ應ご製造セラレダ、輸入セラ付セラルベシ何レカノ年中ニ於テ斯ク交付セラレタルベク且本條約ニ適合スペキ數量ニ於テノミ隨時交レタル際現存スル超過在庫品、政府ニ依り保管セラ右ニ依ラザルトキ、本條約ノ一切/規定が質施セラ

if possible, be utilised for that purpose within the period for which the estimate applies.

In the event, however, of it being impossible to utilise the full amount for that purpose within the period in question, the portion remaining unused at the end of the year shall be deducted from the estimates for that country or territory for the following year.

ARTICLE 9.

If at the moment when all the provisions of the Convention shall have come into force, the then existing stocks of any of the drugs in any country or territory exceeds the amount of the reserve stocks of that drug, which according to the estimates for that country or territory, it is desired to maintain, such excess shall be deducted from the quantity which, during the year, could ordinarily be imported or manufactured as the case may be under the provisions of this Convention.

Alternatively, the excess stocks existing at the moment when all the provisions of the Convention shall have come into force shall be taken possession of by the Government and released from time to time in such quantities only as may be in conformity with the present Convention. Any ルベキ總量ヨリ控除セラルベシ

第十條第四章 禁止及制限

電田林中/

ノ其ノ領域ヨリノ輸出ヲ禁止スベシアセチルモルヒネ」又い其ノ鹽類ヲ 含有スル製剤締約國い「デアセチルモルヒネ」、其ノ鹽類及「デ

古場所への

「指示セラルル官廳」仕向ケラルルコトラ要ストラ得但シ右請求、輸入證明書ラ件ヒ且該證明書含有スル製劑ノ數量ノ該國へノ輸出ヲ許可スルコ其ノ鹽類及「ヂアセチルモルヒネ」又、其ノ鹽類ヲ救用及學術用」必要ナル「ヂアセチルモルとよ」、シザル國ノ政府ヨリ請求ヲ受クルトキ、該國ノ醫ニ、光モ締約國ハ「ヂアセチルモルヒネ」、製造セラ

た際が然

ノ責任ニ於テ分配セラルベシ三 斯ク輸入セラレタル數量・輸入國政府ニ佐り共

第十一條

quantities so released during any year shall be deducted from the total amount to be manufactured or imported as the case may be during that year.

CHAPTER IV .- PROHIBITIONS AND RESTRICTIONS.

ARTICLE 10.

 The High Contracting Parties shall prohibit the export from their territories of diacetylmorphine, its salts, and preparations containing diacetylmorphine, or its salts.

2. Nevertheless, on the receipt of a request from the Government of any country in which diacetylmorphine is not manufactured, any High Contracting Party may authorise the export to that country of such quantities of diacetylmorphine, its salts, and preparations containing diacetylmorphine or its salts, as are necessary for the medical and scientific needs of that country, provided that the request is accompanied by an import certificate and is consigned to the Government Department indicated in the certificate.

 Any quantities so imported shall be distributed by and on the responsibility of the Government of the importing country.

ARTICLE 11.

(領域一於テモ行ハレザルベシシ得以程度一確認セラルルニ非ザレバ何レノ國又解極的又、學術的價値アルコトガ關係政府ノ諸足ザルモノノ取引又、取引ノ為ノ製造、該製產品ノ製産品ニシテ本日階項用又、學術用ニ使用セラレカ」業ノ「エクゴニン、アルカロイド」ヨリ得タル回片ノ「フェナントレン、アルカロイド」又、「コロリニカー

電ヲ超過セザルベク且本條約ノ規定ガ適用セラル電野量と合計並二輸出註文ニ應ズルニ必要ナル数が定醫療用及學術用ノ為ノ右ノ國又、領域ノ内部がご製造ラ許サルル数量、後二視グラルル決定ア産品ニ轉換セラレ保ザルコトラ決定スルニ非ザレラ生ビシメ得ザルカ及、中毒癖ラ生ビシメ得か製右ノ場合ニ 於テハ (政府ニ於ラ右製産品ガ中華辦古ノ場合ニ 於テハ (政府ニ於ラ右製産品ガ中華辦

委員會ニ之ヲ通知スペシ直ニ通告スペク事務総長、他ノ緒約國及聯盟保健エトヲ許ス締約國、其ノ旨ヲ國際聯盟事務總長ニニ 右製産品ノ取引又、取引ノ為ノ製造ヲ開始スル

1. No trade in or manufacture for trade of any product obtained from any of the phenanthrene alkaloids of opium or from the ecgonine alkaloids of the coca leaf, not in use on this day's date for medical or scientific purposes shall take place in any country or territory unless and until it has been ascertained to the satisfaction of the Government concerned that the product in question is of medical or scientific value.

In this case (unless the Government determines that such product is not capable of producing addiction or of conversion into a product capable of producing addiction) the quantities permitted to be manufactured, pending the decision hereinafter referred to, shall not exceed the total of the domestic requirements of the country or territory for medical and scientific needs, and the quantity required for export orders and the provisions of this Convention shall apply.

2. Any High Contracting Party permitting trade in or manufacture for trade of any such product to be commenced shall immediately send a notification to that effect to the Secretary-General of the League of Nations, who shall advise the other High Contracting Parties and the Health Committee of the League. **立扶定** 表员倚人都 開大心保證 石刻遂品。 モノ)ナルカラ決定スペシ 類回又、第二類二掲グラルル薬品ト看像サレ得ツ品ニ轉換シ得ルモノ(其ノ結果トシラ第一類ノ電ルル薬品ト看像サレ得ルモノ)ナルカ又、斯バ薬ルモノ(其ノ結果トシラ第一類ノ電類(二掲グラ金三諮問シタル後右製産品ガ中草獅ラ生ゼラノ得食二路問とない後右製産品ガ中草獅ラ生ゼラノ得

斯夕選任セラレタル二人ニ佐り選任セラルベシ・離盟岡片諸問委員合ニ佐り選任セラレ他ノ一名・問家ノ内一名・開係政府ニ佐り選任セラレ一名・名ノ専門家委員合ニ決定ノ信付託セラルベク右球題・共ノ科學的及技術的方面ノ檢査ヲ高シ得ル三一類ノ電類ロ父・第二類ノ何レニ婦スマナノ問後を得し、「蘇孝君一等の一次の場合ニ於テ・該藥品ガ第日は、は、日、保健委員會ニ於テ右製産品が共レ自體トシテ中回

調け決定人

及第二十七條ニ別グラルル非聯盟國ニ道和スペシ長ニ通告セラルベク事務總長へ之ラ一切ノ聯盟別五 前二號ニ從と到途シタル決定へ國際聯盟事務総

- 3. The Health Committee will thereupon, after consulting the Permanent Committee of the Office international d'Hygiène publique, decide whether the product in question is capable of producing addiction (and is in consequence assimilable to the drugs mentioned in sub-group (a) of Group I), or whether it is convertible into such a drug (and is in consequence assimilable to the drugs mentioned in sub-group (b) of Group I or in Group II).
- 4. In the event of the Health Committee deciding that the product is not itself a drug capable of producing addiction, but is convertible into such a drug, the question whether the drug in question shall fall under sub-group (b) of Group I or under Group II shall be referred for decision to a body or three experts competent to deal with the scientific and technical aspects of the matter, of whom one member shall be selected by the Government concerned, one by the Opium Advisory Committee of the League, and the third by the two members so selected.
- 5. Any decisions arrived at in accordance with the two preceding paragraphs shall be notified to the Secretary-General of the League of Nations, who will communicate it to all the Members of the League and to the non-member States mentioned in Article 27.

関大ル語

ルル適當ナル制度ヲ適用スペシ(第二類ノ何レニ腸スルカニ從と本條約ニ定メラノ通知ヲ受領シタル上右藥品ニ對シ其ノ第一類又ノナリトスルトキハ締約國ハ事務總長ヨリ其ノ旨カ又、中莊癖ヲ生ゼシメ得ル薬品ニ轉換シ得ルモ大 右決定ニシテ右製産品ガ中莊解ヲ生ゼシメ得ル

関上決定人

三佐と變更セラルルコトヲ得来ニ法キ右決定、更ニ得タル經驗ニ盟シ前記手續七 何レカノ締約國ニ佐リ事務總長ニ宛テラルル要

鄉十二樣

本権的で表現の表別の場合の表別の

(レザルベシ域ヨリノ輸出、本條約/規定」従フニ非ザレバ行域ヨリノ輸出、本條約/規定」従フニ非ザレバ行何レカノ藥品ノ締約國ノ領域へノ輸入又、該領

後に記

ヲ控除シテ得タル数量ヲ超過セザルベシリ該年中該國又ハ該領域ニ於テ製造セラルル數量該國又ハ該領域ヨリ輸出セラルル數量トノ合計ヨ」於ケル輸入、第五條ニ定メラルル見額ト該年中二 何レカノ藥品ノ何レカノ國又、領域ヘノ一年間

- 6. If the decisions are to the effect that the product in question is capable of producing addiction or is convertible into a drug capable of producing addiction, the High Contracting Parties will, upon receipt of the communication from the Secretary-General, apply to the drug the appropriate regime laid down in the present Convention according as to whether it falls under Group I or under Group II.
- Any such decision may be revised, in accordance with the foregoing procedure, in the light of further experience, on an application addressed by any High Contracting Party to the Secretary General.

ARTICLE 12.

- No import of any of the drugs into the territories of any High Contracting Party or export from those territories shall take place except in accordance with the provisions of this Convention.
- 2. The imports in any one year into any country or territory of any of the drugs shall not exceed the total of the estimates as defined in Article 5 and of the amount exported from that country or territory during the year, less the amount manufactured in that country or territory in that year.

第五章取締

第十三條



- 株三佐り同條約ノ見定ョり除外セラルルコトアノ製剤ニ適用スペシ但シ「ジュネーヴ」條約第八ン」ノ製劑益二第一類ニ於ケル他ノ薬品ノ一切約第四條ニ包含セラルル「モルセネ」及「コカイ適用スペシ 締約國 パ 又右規定ラ「ジュネーヴ」條合致スル規定) 7第一類ニ於ケル一切ノ藥品ニン物質ニ適用セラルル 同條約ノ規定(ダハ之ニソ 締約國ハ「ジュネーヴ」條約第四條ニ協グラル
 - ト同様ニ取扱フベシラ合有スルモノラ右割合ヲ超エラ含有スルモノヲ右割合ヲ超エラ含有スル製剤ト」以下又ハ「コカイン」つ・1「パーセント」以下液又、稀薄物ニシテ「モルヒネ」つ・1「パーセンケル「モルヒネ」、「コカイン」又い其ノ鹽類ノ溶回、締約國ハ液體又、固體タル無力ノ物質中ニ於

周 上二 締約國(第二類二包含セラレスハ包含セラルル

CHAPTER V .- CONTROL.

ARTICLE 13.

- 1. (a) The High Contracting Parties shall apply to all the drugs in Group I the provisions of the Geneva Convention which are thereby applied to substances, specified in its fourth Article (or provisions in conformity therewith). The High Contracting Parties shall also apply these provisions to preparations made from morphine and cocaine and covered by Article 4 of the Geneva Convention and to all other preparations made from the other drugs in Group I except such preparations as may be exempted from the provisions of the Geneva Convention under its eighth Article.
- (b) The High Contracting Parties shall treat solutions or dilutions of morphine or opcaine or their salts in an inert substance, liquid or solid, which contain 0.2 per cent or less of morphine or 0.1 per cent or less of cocaine in the same way as preparations containing more than these percentages.
 - 2. The High Contracting Parties shall apply

ノ規定(又っ之ニ合致スル規定)ヲ適用スベシコトアルベキ藥品ニ勤シ 左記「ジュネーツ」條約

- 輸出及卸賣ニ關スル限リ該規定、第六條及第七條、規定が右藥品・製造、輸入、
- 付きい之ヲ適用セズ合成物ニシテ普通ノ治療用ニ充テラルルモノニロ 第五章1規定但と右藥品ノ何レカヲ含有スル
- 定担シの第二十二條第一號の、の及中益三第二號へ規
 - ニ迄付セラレ得べク 輸入及輸出ノ統計(毎四半期)代ニ一年毎
 - レカラ合有スル製剤ニ適用セラレザルベシ(第二十二條第一號)日及第二號ハ右藥品ノ何

終十回練

町へ / 連告 政中央条官 三隅 スル常 許可意数結

用セラレザル國父、領域へノ輸出ニ對シ許可證ラベキ藥品ノ本條約及「ジュネーヴ」條約ノ何レモ適一第一類ニ包含セラレ及ハ包含セラルルコトアル

to the drugs which are or may be included in Group II the following provisions of the Geneva Convention (or provisions in conformity therewith):

- (a) The provisions of Articles 6 and 7 in so far as they relate to the manufacture, import, export and wholesale trade in those drugs;
- (b) The provisions of Chapter V, except as regards compounds containing any of these drugs which are adapted to a normal therapeutic use;
- (c) The provisions of paragraphs 1 (b).
 (c) and (c) and paragraph 2 of Article 22, provided:
 - (i) That the statistics of import and and export may be sent annually instead of quarterly, and
 - (ii) That paragraph 1 (b) and paragraph 2 of Article 22 shall not apply to preparations containing any of these drugs.

ARTICLE 14.

1. Any Government which has issued an authorisation for the export of any of the drugs which are or may be included in Group I to any country or territory to which neither this Conven-

モシムベキ敷量ノ輸出ラ許可セザルベシスペキ旨ノ通告ラ後スルトキ、政府へ右超過ヲ生ナセルベキモノトス常設中央委員舎ガ右超過ノ生テ常設中央委員會ヨリ確ムル迄許可證、發給セラティ見積ノ超過ヲ生ゼシメザルコトヲ右政府ニ於以上ナルトキ、行輸出ガ輸入スル國父、領域ニ付員舎三通告スベシ但シ輸出ノ請求ガ五キログラム技術シグル政府へ許可證人發給ヲ前ニ常設中央委

 ○災二在ラズ シ災二輪田ヶ許可セザルベシ但シ左記一場合、比三通告スペク締約國、該年中右ノ國又、領域一動ルルトキ、同委員合、直二右事宜ラ一切ノ締約國タルコトノ判明セル數量トノ和ヲ超過スト認メラ該國又、該領域三付ラノ見荷ノ總量ト輸出セラレ統計「許可セラレタル數量対第五條二定メラルル通告二依リ何レカノ國又、領域二輸出セラレ及、 報告二依リ何レカノ國及、領域二輸出セラレ及、 報告二依リ何レカノ國又、領域二輸出セラレ及、
 ○ tion nor the Geneva Convention applies shall immediately notify the Permanent Central Board of the issue of the authorisation; provided that, if the request for export amounts to 5 kilogrammes or more, the authorisation shall not be issued until the Government has ascertained from the Permanent Central Board that the export will not cause the estimates for the importing country or territory to be exceeded. If the Permanent Central Board sends a notification that such an excess would be caused, the Government will not authorise the export of any amount which would have that effect.

2. If it appears from the import and export returns made to the Permanent Central Board or from the notifications made to the Board in pursuance of the preceding paragraph that the quantity exported or authorised to be exported to any country or territory exceeds the total of the estimates for that country or territory as defined in Article 5, with the addition of the amounts shown to have been exported, the Board shall immediately notify the fact to all the High Contracting Parties, who will not, during the currency of the year in question, authorise any new exports to that country except:

- →提出セラルが場合文へ→福足見就が過剰輸入数量及所要追加數量ニ開
- 場合 ・治療・為缺クベカラザルモノト認ムル例外的 「一輸出國ノ政府ニ於テ輸出ガ人道ノ為父ハ患者

月舎ノ作成の記事中失去

- 年二付左記ラボス表ラ作成スペシ三、常設中央委員會、毎年各國文、各領域ニ闞シ前
 - **~ 各薬品ニ關スが見積**
 - ロ 各葉品ノ市炒数品
 - 、 各藥品 / 製造数量
 - 11 各藥品/與極數品
 - ま 各薬品ノ輸入数量
 - へ 各薬品ノ篠田敷品
 - 使用セラレタル各類品と数量・数量・総由スグニ輸出許可ヲ要セザル製剤ノ製造ニ

示ストキハ右委員合ハ國際解盟事務總長ヲ通ジ右セザリシカ父ハ履行セザリシコトアルベキコトラ右表ガ何レカノ縮約國ノ本條約二依ル義務ヲ履行

- (i) In the event of a supplementary estimate being furnished for that country in respect both of any quantity over-imported and of the additional quantity required; or
- (ii) In exceptional cases where the export in the opinion of the Government of the exporting country is essential in the interests of humanity or for the treatment of the sick.
- 3. The Permanent Central Board shall each year prepare a statement showing, in respect of each country or territory for the pi ceding year:
 - (a) The estimates in respect of each drug;
 - (b) The amount of each drug consumed;
 - (c) The amount of each drug manufactured;
 - (d) The amount of each drug converted;
 - (c) The amount of each drug imported;
 - -(f) The amount of each drug exported;
 - (g) The amount of each drug used for the compounding of preparations, exports of which do not require export authorisations.

If such statement indicates that any High Contracting Party has or may have failed to carry out his obligations under this Convention, the Board shall have the right to ask for explanations, 七麗ニ説グラルル手續の適用セラルベシ合ニの「ジュネーツ」條約第二十四條第二號乃至第締約國ヨリ説明ヲ求ムルノ權利ヲ有スベク此ノ場

キ意見ヲ公表スペシ明~要求ニ關シ其ノ表明セント欲スルコトアルベスハ要求ヒラレクル説明/要領及右ノ説明又ハ説不必要ナリト思考セザル限リ前項ニ従ヒ與ヘラレ右委員會ハ爾後能フ限リ 迷ニ前記/表、委員會ガ

ナル措置ヲ魏ルベシ 公表セラレザルベキコトヲ確保スル為一切ノ必要約國ノ正常ナル商業ヲ阻害スルガ如キ方法ニ佐リ及他、情報ガ投機者ノ行動ヲ容易ナラシブ又、締常設中央委員會ハ本條約ニ依り其ノ受領スル統計

第十五條第六章 行政規定

の方法

偽ニ一切ノ必要ナル立法上又、他ノ措置ヲ執ルベシ締約國、共ノ領域内ニ於ラ本條約ノ規定ヲ宜施スル

through the Secretary-General of the League of Nations, from that High Contracting Party, and the procedure specified in paragraphs 2 to 7 of Article 24 of the Geneva Convention shall apply in any such case.

The Board shall, as soon as possible thereafter, publish the statement above mentioned together with an account, unless it thinks it unnecessary, of any explanations given or required in accordance with the preceding paragraph and any observations which the Board may desire to make in respect of any such explanation or request for an explanation.

The Permanent Central Board shall take all necessary measures to ensure that the statistics and other information which it receives under this Convention shall not be made public in such a manner as to facilitate the operations of speculators or to injure the legitimate commerce of any High Contracting Party.

CHAPTER VI.-ADMINISTRATIVE PROVISIONS.

ARTICLE 15.

The High Contracting Parties shall take all necessary legislative or other measures in order to give effect within their territories to the provisions of this Convention.

別ノ行政機關ヲ改置スベシ締約國ハ既ニ設置シタルニ非ザレバ左記目的ノ為特

- イ 本條約~現定ラ適用スルコト
- ロ 薬品取引ヲ規律シ、監視シ及取締ルコト
- 撲滅運動ラ行フェトル背置ヲ執ルコトニ佐リ中毒癖の 薬品中毒癖ノ蔓延ヲ防止シ及不正取引ヲ禁遏ス

第十六條

張う行うべ際重ナル監

- **今緒約國 5 左記 1 勤 2 聡重ナル監視す行フベン**
- 業者、保有スル原料及既製藥品・數量の 薬品・製造者・轉換又、他・目的・偽各製造
- →数量● 製産セラレタル薬品及、薬品ラ含有スル製剤
- ヨリノ引後 大製産セラレタル薬品及製剤ノ底分称ニ工場

落構ニ關ス関係はノ

11 締約國(市場「情況ラ考慮シタル上事業」經濟

- The High Contracting Parties shall, if they have not already done so, create a special administration for the purpose of:
 - (a) Applying the provisions of the present Convention;
 - (b) Regulating, supervising and controlling the trade in the drugs;
 - (c) Organising the campaign against drug addiction, by taking all useful steps to prevent its development and to suppress the illicit traffic.

ARTICLE 16.

- Each High Contracting Party shall exercise a strict supervision over:
 - (a) The amounts of raw material and manufactured drugs in the possession of each manufacturer for the purpose of the manufacture or conversion of any of the drugs or otherwise;
 - (b) The quantities of the drugs or preparations containing the drugs produced;
 - (c) The disposal of the drugs and preparations so produced with especial reference to deliveries from the factories.
 - 2. No High Contracting Party shall allow

上號ニ説ゲラルル手續へ適用セラルベシ合ニハ「ジュネーツ」條約第二十四條第二號乃至第締約國ヨリ説明ヲ求ムルノ權利ヲ有スベク此ノ場

キ意見ヲ公表スベシ明ノ要求ニ闘シ其ノ表明セント欲スルコトアルベ及ハ要求ニ闘シ其ノ表明セント欲スルコトアルベスハ要求セラレタル説明八要領及右ノ説明又ハ説不必要ナリト思考セザル限リ前項ニ従と與ヘラレ右委員會ハ爾後能フ限リ 迷ニ前記ノ表、委員會ガ

ナル措置ヲ執ルベシ 公表セラレザルベキコトヲ確保スル為一切ノ必要約國ノ正常ナル商業ヲ阻害スルガ如キ方法ニ依リ及他、情報が投機者ノ行動ヲ容易ナラシメ又、締常設中央委員會、本條約ニ依り其ノ受領スル統計

第十五條第 大 章 行政規定

偽ニ一切ノ必要ナル立法上又、他ノ指置ヲ執ルベシ締約國、其ノ領域内ニ於テ本條約ノ規定ヲ宜施スル

through the Secretary-General of the League of Nations, from that High Contracting Party, and the procedure specified in paragraphs 2 to 7 of Article 24 of the Geneva Convention shall apply in any such case.

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別ノ行政機關ヲ設置スペシ羅約國ハ既ニ設置シタルニ非ザレバ左記目的ノ為特

- イ 本條約ノ規定ヲ適用スルコト
- ロ 薬品取引ヲ規律シ、監視シ及取締ルコト
- 撲滅運動ヲ行フコトル為一切ノ右用ナル措置ヲ執ルコトニ佐リ中毒癖() 薬品中毒癖ノ蔓延ヲ防止シ及不正取引ヲ禁遏ス

第十六條

観り行う。

- 各緒約國、左記ニ對シ嚴重ナル監視ヲ行フベシ
- 業者、保有スル原料及既製藥品ノ數量(薬品・製造者・轉換又、他・目的)為各製造
- 収量製産セラレタル薬品及、薬品ラ含有スル製剤
- ヨリノ引波 右製産セラレクル薬品及製剤ノ底分符ニ工場

落陣二隅ス関連原料人

11 締約國、市場、情况ラ考慮シタル上事業・經濟

- The High Contracting Parties shall, if they have not already done so, create a special administration for the purpose of:
 - (a) Applying the provisions of the present Convention;
 - (b) Regulating, supervising and controlling the trade in the drugs;
 - (c) Organising the campaign against drug addiction, by taking all useful steps to prevent its development and to suppress the illicit traffic.

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- Each High Contracting Party shall exercise a strict supervision over:
 - (a) The amounts of raw material and manufactured drugs in the possession of each manufacturer for the purpose of the manufacture or conversion of any of the drugs or otherwise;
 - (b) The quantities of the drugs or preparations containing the drugs produced;
 - (c) The disposal of the drugs and preparations so produced with especial reference to deliveries from the factories.
 - 2. No High Contracting Party shall allow

ル取締

「供谷量ヲ超過セザルベシニ於テモ蓄積セラルルコトアルベキ總量、一年分ト器ムルトキ、此ノ限ニ在ラザルモ如何ナル場合ノ後例外的事情ニ依リ追加數量ノ蓄積ヲ正當ナリル數量ヲ超過セザルベシ但シ政府ガ充分ナル調査ル数量モ右製造業者が確後ノ大月間ノ製造ニ要ス者ノ干許二保有セラルル原料ノ如何ナル時ニ於ケ治競者・手許ニ於ケル蓄積ヲ許サザルベシ製造業

第十七条

現はスペキ関語来者

記被スル四中期報告ヲ提出スルコトヲ要求スベン各締約國い自國ノ領域内ノ各製造業者ニ對シ左記ヲ

作が充分ナット認ムが條件ノ下ニ決定セラレタルン」ノ割合ニシテ政府ノ規定スが方法ニ佐り且政権と得が「モルヒネ」、「コカイン」又ハ「エクゴニ量ヲ報告スルニ當リ之ニ合有セラレ又ハ之ヨリ製産品ノ數量製造業者ハ右受入レタル原料ノ数數量就ニ右各物質ヨリ製産セラレタル薬品又ハ他付替認造業者が工場ニ受入レタル原料及各藥品ノ

the accumulation in the possession of any manufacturer of quantities of raw materials in excess of those required for the economic conduct of business, having regard to the prevailing market conditions. The amounts of raw material in the possession of any manufacturer at any one time shall not exceed the amounts required by that manufacturer for manufacture during the ensuing six months, unless the Government, after due investigation, considers that exceptional conditions warrant the accumulation of additional amounts, but in no case shall the total quantities which may be accumulated exceed one year's supply.

ARTICLE 17.

Each High Contracting Party shall require each manufacturer within his territories to submit quarterly reports stating:

(a) The amount of raw materials and of each of the drugs received into the factory by such manufacturer and the quantities of the drugs, or any other products whatever, produced from each of these substances. In reporting the amounts of raw materials so received, the manufacturer shall state the proportion of morphine, cocaine or ecgonine contained in or producible therefrom as deter-

モノヲ記載スペシ

提出スペキ四段集合ノ

期中三庭分セラレタル数量の右原料又、之ヨリ製造セラレタル製産品ノ四年

3 四 期末二於ケル殘存在庫數量

コトラ要求スペシ 右藥品ノ數点ヲ記載スル報告ヲ年末ニ於テ提出スル各年中輸出文、輸入セラレタルモノニ合介セラルルニ開シ、輸出文、輸入ニ許可ヲ要セザル製劑ニシテ各締約國、自國ノ領域内ノ各組資業者ニ對シ各藥品

第十八條

度な漢品~

一充テラルベキコトラ約ス一切/場合二於ティデア性ナキ物質二轉換セラルルカダへ溶療用者へ學術用作二依り父へ近ノ収縮ノ下二原薬セラルルカ、麻酔動ノ為二必要ナラザルニギリタルトキへ右藥品へ政何レカノ藥品ガ网ノ機關ニ依ル司法子額又、他ノ行各緒約例へ不正取引二於テ其ノ押收シタル第一類ノ各緒約例。不正取引二於テ其ノ押收シタル第一類ノ

mined by a method prescribed by the Government and under conditions considered satisfactory by the Government;

(b) The quantities of either the raw material or the products manufactured therefrom which were disposed of during the

quarter;
(c) The quantities remaining in stock at the end of the quarter.

Each High Contracting Party shall require each wholesaler within his territories to make at the close of each year a report stating, in respect of each of the drugs, the amount of that drug contained in preparations, exported or imported during the year, for the export or import of which authorisations are not required.

ARTICLE 18.

Each High Contracting Party undertakes that any of the drugs in Group I which are seized by him in the illicit traffic shall be destroyed or converted into non-narcotic substances or appropriated for medical or scientific use, either by the Government or under its control, when these are no longer required for judicial proceedings or other action on the part

ベシセチルモルとネ」「関東セラルルカダハ轉換セラル

第十九條

上記載スパンプルル

又國内法令ニ於テ定メラルル藥品名ヲ示スベシ行牟ヲテスペキコトヲ要求スペシ行「Vァテル」ニハスルガ為ニ使用セラルル「Vァテル」ニハ該藥品ノ含縮約國ハ何レカノ藥品又ハ之ヲ含有スル製劑ヲ販賣

第七章一般規定

総二十条



参え、商祉ノ名及気所ラ示スペシ製造セラレス、耐換セラルル薬品並ニ許可セラルモ目的トスルサ、右製造文、轉換ノ間結スル日、、轉換方内部指要ノ為ノミナリセス、更二輸出ラト欲スルトキ、网際聯盟事務総長ニ通告シ製造文

「有宜施ノ際者、爾後右製造者、轉換り許可セン於子何レカノ藥品が製造中者、轉換中ナルトキ又今緒約國、本條約宜施ノ際共ノ領域ノ何レカニ

of the authorities of the State. In all cases diacetylmorphine shall either be destroyed or converted.

ARTICLE 19.

The High Contracting Parties will require that the labels under which any of the drugs, or preparations containing those drugs, are offered for sale, shall show the percentage of the drugs. These labels shall also indicate the name of the drugs as provided for in the national legislation.

CHAPTER VII.—GENERAL PROVISIONS.

ARTICLE 20.

1. Every High Contracting Party in any of whose territories any of the drugs is being manufactured or converted, at the time when this Convention comes into force, or in which he proposes either at that time or subsequently to authorise such manufacture or conversion, shall notify the Secretary-General of the League of Nations indicating whether the manufacture or conversion is for domestic needs only or also for export, the date on which such manufacture or conversion will begin, and the drugs to be manu-

- 係商社ノ名及宛所ラ示スペシムペキ場所及日ヲ示シ且右藥品並ニ關係者又、關其ノ旨ヲ道告シ右製造又、轉換ガ止ミタル又、止ノ領域ニ於テ止ム場合ニ、該締約國、事務總長ニニ、何レカノ藥品ノ製造又、轉換ガ何レカノ締約國
- 縮約國ニ道知セラルベシ三二 本條二依り供與セラルル情報(事務總長二依り

继11十1条

成セラルル様式ニ従と事務鵜長ニ登付スペシ及他」危险藥品ノ取引ニ關スル諸問委員會ニ依り作其ノ領城ニ於ケル本條約ノ運用ニ關スル年限ヲ阿庁則ヲ國際聯盟事務總長ヲ通ジテ相互ニ通知スペク且縮約國ハ本條約ヲ宜施スル為公布セラルル法令及規

総11十112

factured or converted as well as the names and addresses of persons or firms authorised.

- 2. In the event of the manufacture or convension of any of the drugs ceasing in the territory of any High Contracting Party, he shall notify the Secretary-General to that effect, indicating the place and date at which such manufacture or conversion has ceased or will cease and specifying the drugs affected, as well as the names and addresses of persons or firms concerned.
- The information furnished under this Article shall be communicated by the Secretary-General to the High Contracting Parties.

ARTICLE 21.

The High Contracting Parties shall communicate to one another through the Secretary-General of the League of Nations the laws and regulations promulgated in order to give effect to the present Convention, and shall forward to the Secretary-General an annual report on the working of the Convention in their territories, in accordance with a form drawn up by the Advisory Committee on Traffic in Opium and Other Dangerous Drugs.

ARTICLE 22.

建田年大統計人

ラ包含セシムベシスル為製造業者及卸賣商が使用シタル各藥品・数量ヲ問ハズ輸出スルニ輸出許可予要セザル製剤ヲ製造ハ内部消費ノ為ノモノタルト輸出ノ為ノモノタルト締約國ハ共ノ常設中央委員會ニ提出スル年次統計ニ

~概要ラ右統計ニ包含セシムベツが衝勢の右続計=包含セシムベツが観光を手上後ニ從ヒ製造業者/作成スル報告

解11十111森

開スル西部

速ニ相互ニ道知スペシモノノ詳細事項ヲ國際聯盟事務總長ヲ通ジ能フ限リタル方法ニ闢スル資料ニ徴シ重要ナルコトアルベキレタル藥品ノ出所者、不正取引者ニ依り使用セラレ該藥品ノ數量ニ徴シ及、不正取引ノ目的ヲ以テ得ラ締約國、北ノ後見シタル不正取引ノ各場合ニシテ嘗

右詳細事項ニ、能フ限リ左記ヲ示スペシ

- **~ 常該薬品 / 種類及数量**
- p 藥品 / 原産地、記號及「ソッテル」

The High Contracting Parties shall include in the annual statistics furnished by them to the Permanent Central Board the amounts of any of the drugs used by manufacturers and wholesalers for the compounding of preparations whether for domestic consumption or for export for the export of which export authorisations are not required.

The High Contracting Parties shall also include a summary of the returns made by the manufacturers in pursuance of Article 17.

ARTICLE 23.

The High Contracting Parties will communicate to each other, through the Secretary-General of the League of Nations, as soon as possible, particulars of each case of illicit traffic discovered by them which may be of importance either because of the quantities involved or because of the light thrown on the sources from which drugs are obtained for the illicit traffic or the methods employed by illicit traffickers.

The particulars given shall indicate as far as

- (a) The kind and quantity of drugs
- involved;
 (b) The origin of the drugs, their marks and labels;

- ハ 薬品ガ不正取引ニ韓向セラレタル場所
- 及宛所人人名、荷弦方法並ニ判明セルトキ、荷受人ノ名、古弦方法並ニ判明セルトキ、荷受人ノ名」、秦品ノ後迩地、迅弦人若、迅迩取扱人又、荷弦
- 品 「 街途セラレタル船舶アルトキ (共)名称 第 注 と を 総 発 注 一 後 総 発 注 (後 転 方) 女) 使 用 セラレタル 方 法 及 総 路 道 二 薬
- ノ執りタル行動及科セラレタル刑罰(開係者特ニ許可又の免許ラ行スル者ニ闘シ政府)
- ト 不正取引ノ禁遏ニ査スルコトアルベキ他ノ情報

終二十回森

ラルル稲約岡問ノ開係ニ於テ此等ノ條約ヲ補足スベ五年ノ「ジュネーツ」條約ノ少クトモ 一ニ佐り 拘束セ本條約ハ千九百十二年ノ「ヘーグ」條約及千九百二十

郑11十五株

理又へ適用本様約~解

本條約ノ解釋文、適用ニ闢シ締約國問ニ何レカノ粉

- (c) The points at which the drugs were diverted into the illicit traffic;
- (d) The place from which the drugs were despatched, and the names of shipping or forwarding agents or consignors; the methods of consignment and the name and address of consignees, if known;
- (c) The methods and routes used by smugglers and names of ships, if any, in which the drugs have been shipped;
- (f) The action taken by the Government in regard to the persons involved, particularly those possessing authorisations or licences and the penalties imposed;
- (g) Any other information which would assist in the suppression of illicit traffic.

ARTICLE 24.

The present Convention shall supplement the Hague Convention of 1912 and the Geneva Convention of 1925 in the relations between the High Contracting Parties bound by at least one of these latter Conventions.

ARTICLE 25.

If there should arise between the High Con-

事に関スル紛

二開スル協定二從と解決セラルベシキ(右紛字(當事國問ニ質施セラルル國際紛爭解決事務生の外次國際紛爭解決事發生の外交手段二依り滿足二解決と得ラレザルト

採二十六茶

開スル領田衛民地等に

部父(何レカニ腸と何等ノ養務ラモ負(ギルコトラ外領主父(宗主権方(委任統治ノ下ニ在ル地域ノ全権約國(本條約ヲ受能スルモ共ノ殖民地、保護領、海

tracting Parties a dispute of any kind relating to the interpretation or application of the present Convention and if such dispute cannot be satisfactorily settled by diplomacy, it shall be settled in accordance with any applicable agreements in force between the Parties providing for the settlement of international disputes.

In case there is no such agreement in force between the Parties, the dispute shall be referred to arbitration or judicial settlement. In the absence of agreement on the choice of another tribunal, the dispute shall, at the request of any one of the Parties, be referred to the Permanent Court of International Justice, if all the Parties to the dispute are Parties to the Protocol of December 16th, 1920, relating to the Statute of that Court, and, if any of the Parties to the dispute is not a Party to the Protocol of December 16th, 1920, to an arbitral tribunal constituted in accordance with the Hague Convention of October 18th, 1907, for the Pacific Settlement of International Disputes.

ARTICLE 26.

Any High Contracting Party may, at the time of signature, ratification or accession, declare that, in accepting the present Convention, he does not assume any obligation in respect of all or any

レザルベシ佐り本体約ハ右宣言中二掲グラルル地域ニ適用セラ保名、批准文、加入ノ際官言スルコトラ得ベク之ニ

ルル一切ノ地域ニ盗用セラルベシニ加入スル國ノ場合ニ於ケルト同様右道知ニ以グラルコトラ得ベク之ニ依り本條約ハ之ヲ批准シ又ハ之スル言ヲ解後何時ニテモ國際聯盟事務總長ニ道知スノ各部又ハ何レカニ本條約ヲ適用セラルルコトラ欲締約國ハ前項ニ依ル官言ノ目的ト為リタル共ノ地域

城三適用ナキニ至ルベシル腹塞ノ道告ニ於ケルト同様右官言ニ招ゲラルル連スルコトヲ得ベク之ニ依り本條約ハ同條ノ現定ニ依條ニ投ゲラルル五年ノ即問ノ滿了後何時ニテェ官言本條約ノ適用ナキニ至ルコトヲ欲スル旨ヲ第三十二本餘約ノ適用ナキニ在ル連城ノ各部父ハ何レカニ對シ紅春紀治ノ中ニ在如迪域ノ各部父ハ何レカニ對シ羅約例ハ北ハ所民地、保護領、海外領土父ハ宗主權方

事務總長、本條三依り受領シタル一切ノ宜言及通知

of his colonies, protectorates and overseas territories or territories under suzerainty or mandate, and the present Convention shall not apply to any territories named ln such declaration.

Any High Contracting Party may give notice to the Secretary-General of the League of Nations at any time subsequently that he desires that the Convention shall apply to all or any of his territories which have been made the subject of a declaration under the preceding paragraph, and the Convention shall apply to all the territories named in such notice in the same manner as in the case of a country ratifying or acceding to the Convention.

Any High Contracting Party 100, at any time after the expiration of the five-years period mentioned in Article 32, declare that he desires that the present Convention shall cease to apply to all or any of his colonies, protectorates and overseas territories or territories under suzerainty or mandate, and the Convention shall cease to apply to the territories named in such declaration as if it were a denunciation under the provisions of Article 32.

The Secretary-General shall communicate to

ア一切ノ韓盟國及第二十七後二极ゲラルル非諸盟國 三道様スパシ

経11十九茶

條約、能蘭西語及英吉利語ノ本文ヲ以テ共ニ正文 トッ本ロノ日附ラ行スペク且國際聯盟ノ聯盟國又、 本候約ヲ作成シタル合議二代表者ヲ出シタル非聯盟 岡若、岡際韓盟理事合ガ本條約ノ謄本ヲ署名ノ為送 性シタル非難盟國/男名/為于九百三十一年十二月 三十一日空間を置さかべる

第二十八條

甘

本條約、批准セラルベン批准書、國際聯盟事務總長 三笠付セラルベク事務総長、之が受領ラ一切ノ聯盟 國及前條二极グラルル非韓盟國ニ通告スペシ

第二十九條

十九百三十二年一日一日以後国際韓盟ノ韓盟國文(

all the Members of the League and to the nonmember States mentioned in Article 27, all declarations and notices received in virtue of this Article.

ARTICLE 27.

The present Convention, of which the French and English texts shall both be authoritative, shall bear this day's date, and shall, until December 31st, 1931, be open for signature on behalf of any Member of the League of Nations, or of any nonmember State which was represented at the Conference which drew up this Convention, or to which the Council of the League of Nations shall have communicated a copy of the Convention for this purpose.

ARTICLE 28.

The present Convention shall be ratified. The instruments of ratification shall be transmitted to the Secretary-General of the League of Nations, who shall notify their receipt to all Members of the League and to the non-member States referred to in the preceding Article.

· ARTICLE 29.

As from January 1st, 1932, the present Con-

第二十七條二叔グラルル非解盟國、本條約二加入ス ルコトラ得

加入告へ國際聯盟事務總長ニ迄付セラルベク事務總 長いたが受領ラー切り韓盟國及第二十七條三极ゲラ ガル非韓盟國ニ道告アベッ

郑川十紫

54

本條約、國際聯盟事務總長が在ノ諸國ノ內四國ヲ含 ムニ十五ノ网際韓盟ノ韓盟國文、非韓盟國ノ批准文 い加入ラ受削シタル後九十日ニシラ宣龍セラルベシ

俳單内図、図逸図、「グレート、ブリテン」及北部「ア イドランド」縣今下風、二本風、和田園、龍西風、「下 ルコ」国及「アメリカ」合衆国

尤丰第二條乃至第五條以外,本條約,規定,第二條 乃至第五條三從と提出セラルル見積ノ關スル最初ノ 年ノ一月一日ヨリノミ適用セラルベキモノトス

经川十一条

vention may be acceded to on behalf of any Member of the League of Nations or any nonmember State mentioned in Article 27.

The instruments of accession shall be transmitted to the Secretary-General of the League of Nations, who shall notify their receipt to all the Members of the League and to the nonmember States mentioned in that Article.

ARTICLE 30.

The present Convention shall come into force ninety days after the Secretary-General of the League of Nations has received the ratifications or accessions of twenty-five Members of the League of Nations or non-member States, including any four of the following:

France, Germany, United Kingdom of Great Britain and Northern Ireland, Japan, Netherlands, Switzerland, Turkey, and the United States of America.

Provided always that the provisions of the, Convention other than Articles 2 to 5 shall only be applicable from the first of January in the first year in respect of which estimates are furnished in conformity with Articles 2 to 5.

ARTICLE 31.

本條約ノ賞施ノ日ノ後受領セラルル批准又、加入: 國際聯盟事務總長ガ之ヲ受領シタル日ヨリ九十日ノ 期間ノ流了ノ時ヨリ效力ヲ生ズベシ

海川十川海

本條約ノ宜施ノ日ヨリ五年ノ期問滿了後ニ於テハ本 條約、國際聯盟事務總長ニ常託セラルル書面ニ依り 廢棄セラルルコトラ得廢棄、何レカノ年ノ七月一日 以前二年務總長二依り受領セラルルトキの翌年ノー 月二日二效力ラ生ズベク七月一日後二受領セラルル トキハ翌年ノ七月一日以前ニ受領セラレタルモノト シラ效力ラ生ズベン各検薬ハ之ラ容託シタル離盟國 又、非辭盟國ニ對シテノミ有效ナルベシ

事務總長、受領シタル廢棄ラ一切ノ聯盟國及第二十 七條二极グラルル非韓盟國ニ通告スペシ

Ratifications or accessions received after the date of the coming into force of this Convention shall take effect as from the expiration of the period of ninety days from the date of their receipt by the Secretary-General of the League of Nations.

ARTICLE 32

After the expiration of five years from the date of the coming into force of this Convention, the Convention may be denounced by an instrument in writing, deposited with the Secretary-General of the League of Nations. The denunciation, if received by the Secretary-General on or before the first day of July in any year, shall take effect on the first day of January in the succeeding year, and, if received after the first day of July, shall take effect as if it had been received on or before the first day of July in the succeeding year. Each denunciation shall operate only as regards the Member of the League or non-member State on whose behalf it has been deposited.

The Secretary-General shall notify all the Members of the League and the non-member States mentioned in Article 27 of any denunciations received.

同時又、順次ノ廢棄ノ結果トシラ本條約ノ拘束ヲ受 クル聯盟國及非聯盟國ノ敷ガニ十五未滿ニ滅少スル トキハ本條約の右廢棄ノ最後ノモノガ本條ノ規定ニ 従と效力ヲ生ズベキ日ヨリ賞施セラレザルニ至ルベ

経111十111条

検約~改正

本條約ノ改正ノ要求、本條約ノ拘束ヲ受クル國際辦 盟ノ聯盟國叉、非聯盟國ニ佐リ國際聯盟事務總長ニ 宛テタル通知ヲ以テ何時ニテモ為サルルコトヲ得右 通知、事務總長二佐り本條約ノ拘束ヲ受クル他ノ國 際聯盟ノ聯盟國又、非聯盟國ニ通報セラルベク其ノ 三分ノ一以上二佐り承認セラルルトキ、締約國、本 條約ノ改正ノ為合合スルコトラ約ス

経川十瓦森

本條約(其ノ貨施ノ日ニ於テ國際聯盟事務總長ニ依

If, as a result of simultaneous or successive denunciations, the number of Members of the League and non-member States bound by the present Convention is reduced to less than twentyfive, the Convention shall cease to be in force as from the date on which the last of such denunciations shall take effect in accordance with the provisions of this Article.

ARTICLE 33.

A request for the revision of the present Convention may at any time be made by any Member of the League of Nations or non-member State bound by this Convention by means of a notice addressed to the Secretary-General of the League of Nations. Such notice shall be communicated by the Secretary-General to the other Members of the League of Nations or non-member States bound by this Convention, and, if endorsed by not less than one-third of them, the High Contracting Parties agree to meet for the purpose of revising the Convention.

ARTICLE 34.

The present-Convention shall be registered

り登録セラルベシ

米 ヌ 行證據トシテ前記を構奏員へ本條約二署名セリ

逐制置

「アメリカ」合案図 ドクトル、カーレル フライ(ル、フォン、ラインバーバン

> サンボーン、ヤング シャがター、バーイス、トンップウェインリー、ジェー、アンスリンガー

其ノ一切ノ誘導體及合成ノ方法ニ佐り製産① 「アメリカ」合衆國政府(阿片、「ヨカ」薬、

by the Secretary-General of the League of Nations on the day of its entry into force.

IN FAITH WHEREOF the above-mentioned Plenipotentiaries have signed the present Convention

Done at Geneva the thirteenth day of July, one thousand nine hundred and thirty-one, in a single copy, which shall remain deposited in the archives of the Sccretariat of the League of Nations, and certified true copies of which shall be delivered to all the Members of the League and to the non-member States referred to in Article 27.

GERMANY

ALLEMAGNE

Freiherr von Rheinbaben

UNITED STATES
OF AMERICA

ETATS-UNIS D'AMÉRIQUE

John K. CALDWELL
Harry J. Anslinger
Walter Lewis Treadway.
Sanborn Young.

(1) The Government of the United States of America reserves the right to impose, for purpose of internal control and control of import into and export from territory under

スルノ権利ヲ留保ス 緒ノ為條約ノ規定ヨリモ嚴重ナル措置ヲ課領域へノ輸入又六其ノ領域ヨリノ輸出ノ取 セラルル同様ノ物質ノ國内取締ノ為及其ノ

- 前提條件ト為シ得が措置ヲ課スルノ権利ヲノ提出ヲ其ノ領域内ノ通過ノ許可ヲ與フル、提出ヲ其ノ領域内ノ通過ノ許可ヲ與フル過ヲ取締ル為仕向國ノ疫給セル輸入許可證製産セラルル同様ノ物質ノ其ノ領域内ノ通薬、其ノ一切ノ誘導體及合成ノ方法ニ依リ□「アメリカ」合衆國政府ハ生阿片、「コカ」
- ラ約スルコト實行不可能ナリト認と 設中央阿片委員舎ニ右統計ヲ恣付スルコトスル三月ノ期間ノ終了後六十日未滿内ニ常三「アメリカ」合衆國政府、輸出入統計ノ關
- コトラ約スルコト宣行不可能ナリト認よ及い輸入セラレタル藥品ノ数显ヲ別ニ示ス問「アメリカ」合衆國政府の政府用ノ為購入
- リカ」 合衆國ノ為ニ麻藥ノ製造制限及分配玉 「アメリカ」合衆國全権委員、本日「アメ

its jurisdiction, of opium, coca leaves, all of their derivatives and similar substances produced by synthetic process, measures stricter than the provisions of the Convention.

- (2) The Government of the United States of America reserves the right to impose, for purposes of controlling transit through its territories of raw opium, coca leaves, all of their derivatives and similar substances produced by synthetic process, measures by which the production of an import permit issued by the country of destination may be made a condition precedent to the granting of permission for transit through its territory.
- (3) The Government of the United States of America finds it impracticable to undertake to send statistics of import and export to the Permanent Central Opium Board short of sixty days after the close of the three-months' period to which such statistics refer.
- (4) The Government of the United States of America finds it impracticable to undertake to state separately amounts of drugs purchased or imported for Government purposes.
- (5) Plempotentiaries of the United States of America formally declare that the signing of the Convention for Limiting the Manufacture and Regulating the Distribution

チェ非サルコトラ正式 | 官言ス予承認スルモノナリトノ意義ニ解セラルベ来國ガ右組織又、質體ラ右ノ國ノ政府トシ依り未ダ永認セラレザル限リ「アメリカ」合政府ナリトシテ「アメリカ」合衆國政府ニニ界名シ又、加入スル組織又、質體ガ一國取締ニ關スル條約二界名スルコト、該條約

グプリュー、エル、ティーエイチ、ジェー、エージェー、シェー、ケー、シー

of Narcotic Drugs by them on the part of the United States of America on this date is not to be construed to mean that the Government of the United States of America recognises a regime or entity which signs or accedes to the Convention as the Government of a country when that regime or entity is not recognised by the Government of the United States of America as the Government of that country.

(6) The plenipotentiaries of the United States of America further declare that the participation of the United States of America in the Convention for limiting the Manufacture of and regulating the Distribution of Narcotic Drugs, signed on this date, does not involve any contractual obligation on the part of the United States of America to a country represented by a regime or entity which the Government of the United States of America does not recognise as the Government of that country until such country has a Government recognised by the Government of the United States of America.

J. K. C H. J. A. W. L. T.

エス、ワイ

フェルナンド、ベレス政府ノ承認ヲ條件トス「アルゼンティン」共和國

製出生成

ドクトル・グがノー・シュガッチー・プット・グル

白耳線図

ドクトガ、エフ、ド、ミットネーグ

「キッシュア」図

エメノクィットーグ

「アレッグ」図

ラウル、ド、リオ、ブランコ

ノー切り部分 益ニ國際輸出ノ側個ノ離盟國ニ非ザル英帝國「グレート、ブリンテン」及北部「アイルランド」

トチロインドフタンソロロ

「キナキ」

S. Y.

ARGENTINE REPUBLIC RÉPUBLIQUE ARGENTINE

* Ad referendum.

Fernando Perez

AUSTRIA

AUTRICHE E. Pflügl

D 0

D' Bruno SCHULTZ

Belgique

D' F. DE MYTTENAERE

Bolvie

M. CUELLAR

Bresil

Raul do Rio Branco

GREAT BRITAIN AND

NOTHERN IRELAND

and all parts of the British Empire which are not separate Members of the League of Nations.

GRANDE-BRETAGNE ET

IRLANDE DU NORD

de l'Empire britannique non Membres séparés de la Société des Nations

Malcolm DELEVINGNE

CANADA

BELGIUM

BOLIVIA

BRAZIL

CANADA

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ダブシュートナートリデル	W. A. RIDDELL
亞 赵	India
アーグ、カー、ベルリカー	R. P. Paranjpye
[45]屋	Сніга
エンタケ、ホータ、ガ・ルド、ヴェー	Enrique J. Gajardo V.
「ロスターカ」図	COSTA RICA COSTA-RICA
ガナジアト、フィグェファ、ロラ	Viriato Figueredo Lora.
「井川、」「屋	CUBA CUBA
(一つがいかか)	G. DE BLANCK
アクトラストングンメンドス	Dr B. PRIMELLES
	Denmark Danemark
一株図	Gustav Rasmussen
グ K W ーレ、レ K イッカッ	FREE CITY OF DANZIG VILLE LIBRE DE DANTZIG
Lダンチッカ ¹ 年年治	F. SOKAL
エレノンセグ	
「ドミュカ」共和國	DOMINICAN REPUBLIC RÉPUBLIQUE DOMINICAINE
カー、アッヤグトツ	Ch. Ackermann
「モジント」図	Едүрт
アナー・ダンシュー・レン・カグ	T. W. Russell
西班子國	Spain Espagne

フリオ、カサレス

「エティオピア」図

エントット公、伯爵ラガルド

高麗因國

ル郊民地、保護領及委任統治地域ニ闘シーニ提出と得かす否ヤニ付其ノ権力ノ下ニ在統計ヲ酸ニ付與セラレタル期間内ニ規則的佛蘭西國政府、第十三條ニ掲グラルル四半期

切〉留保了為ス

シャーノグラント

华嚴國

アーグ、ラントエグ

「グトルトル」図

ガイス、マガティネス、モント

「ヘデァーズ」及「ネデド」開立二属地

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伊太利國

カサトッシャル、ステファノ

Julio CASARES

ABYSSINIA

Етнюрів

Cte LAGARDE DUC d'ENTOTTO

FRANCE

FRANCE

Le Gouvernement français fait toutes ses réserves en ce qui concerne les colonies, protectorats et pays sous mandat dépendant de son autorité, sur la possibilité de produire régulièrement dans le délai strictement imparti les statistiques trimestrielles

visées par l'article 13.
G. Bourgois

GRÈCE

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GUATÉMALA

Luis MARTÍNEZ MONT.

R. RAPHAËL

HEJAZ, NEJD AND DEPENDENCIES HEDJAZ, NEDJED ET DÉPENDANCES

HAFIZ WAHBA

ITALY

GREECE

GUATEMALA

ITALIE

CAVAZZONI Stefano

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縣大十九 麻藥條約 麻藥ノ製造御職以分善即鄉ニ購スス領事		
	ダブリュー、エー、リデルシー・エイチ、エル、シァーァン	C. H. L. SHARMAN W. A. RIDDELL
	色製	INDIA
	アーグ、ガー、ベシュガー	R. P. Paranjpye
	「十二国	CHILE CHILI
	エンリケ、ホータ、ガイルド、ヴェー	Enrique J. Gajardo V.
	「カベダース」図	Costa Rica Costa-Rica
	ガイリアトハフィグェファ、ロラ	Viriato Figueredo Lora.
	「井川大」屋	Сива
	くー、アンシック	G. DE BLANCK
	テクトラングーングンメンドス	Dr B. Primelles
	卜 茶図	DENMARK DANEMARK
	グスターレッシスペッセン	Gustav Rasmussen
	「ダンチッと」年由市	FREE CITY OF DANZIG VILLE LIBRE DE DANTZIG
	サンシャグ	F. Sokal
	「ドミニカニ共和図	DOMINICAN REPUBLIC RÉPUBLIQUE DOMINICAINE
	サートアッケグトン	Ch. Ackermann
,	「月沙人士」屋	Едүрт Едүрте
	ティー・ダブリュー・フッセル	T. W. RUSSELL
	西班子園 ・	Spain Espagne
	THE PAYMENT .	

フリオ、カサレス

「エティオピア」図

エントット公、伯酔ラガルド

年据和

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シャーノグラント

华職國

アーグ、ロントイン

「グァテトル」図

ガイス、マグティネス、モント

「ヘデァーズ」及「ネデド」関立二関地

イハナドハード

伊大利國

カヴァッシャニ、ステファノ

Julio CASARES

ABYSSINIA

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G. Bourgors

GREECE

GRÈCE

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HEJAZ, NEJD AND DEPENDENCIES

HEDJAZ, NEDJED ET DÉPENDANCES

HAFIZ WAHBA

ITALY

ITALIE

CAVAZZONI Stefano

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PAYS-BAS

PERSE

POLOGNE

PORTUGAL

ROUMANIE

SIAM

SUÈDE

SAINT-MARIN

日本國 澤田館藏 大蓬茂雄 「マイント」図 アクトグ、アー、ソッチィーグ 「リベリア」共和國上院ノ批准ヲ留保ス 「リストニト」図 サウニウス 「ラクカソンラグ」図 セーンジェー、ヴェがメーグ 「メキショ」図 エセマアティネス、ディアルヴァ 「キナロ」図 カー、アンチ 「ベナト」図 ドクトル、エルネスト、ホファン し、シグァイ」図

竹蘭國

JAPON JAPAN S. SAWADA S. OHDACHI LIBÉRIA LIBERIA Dr A. SOTTILE Sous réserve de ratification du Sénat de la République de Libéria. LITHUANIE LITHUANIA ZAUNIUS. LUXEMBOURG LUXEMBURG Ch. G. VERMAURE MEXIQUE Mexico S. MARTÍNEZ DE ALVA Monaco MONACO C. HENTSCH. PANAMA PANAMA Dr Ernesto Hoffmann. PARAGUAY PARAGUAY R. V. CABALLERO DE BEDOYA

v. Wettum

A. SEPAHBODY

Снорźко

Augusto DE VASCONCELLOS

A. M. FERRAZ DE ANDRADE

C. ANTONIADE

THE NETHERLANDS

PERSIA

POLAND

PORTUGAL

ROUMANIA

SAN MARINO

遥羅國有害智情性藥品法、或難二於テ「ジョ波性國 選羅國 フェルリ、シァルル、エミール 「サン、マリノ」國 シー、アントニアーデ 「ルーマニア」國 アー、エメ、フェルラス、ディア・デラーデ 「ボルトガル」國 エー、セン・二國 ボデコ 「ボーランド」國 エー、セバーボディ

DAMRAS

As our Harmful Habit-forming Drugs
Law goes beyond the provisions of
the Geneva Convention and the present Convention on certain points,
my Government reserves the right to
apply our existing law.

SWEDEN

ロー、イー、シェベトトン

ラ留保ス

瑶丰匮

以ラ我政府が共ノ現行法ヲ適用スルノ権利ネーヴ」條約及本條約ヨリモ 一層霰ナルラ